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Europäisches Patentamt
European Patent Office
Office européen des brevets

(11)

EP 1 092 435 A1

(12)

EUROPEAN PATENT APPLICATION

published in accordance with Art. 158(3) EPC

(43) Date of publication:

18.04.2001 Bulletin 2001/16

(51) Int. Cl.⁷:**A61K 31/505**

(21) Application number: 99926903.8

(86) International application number:

PCT/JP99/03583

(22) Date of filing: 02.07.1999

(87) International publication number:

WO 00/01388 (13.01.2000 Gazette 2000/02)

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE

(30) Priority: 02.07.1998 JP 18770598

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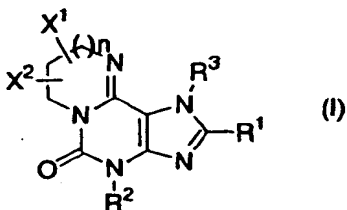
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(54) REMEDIES FOR DIABETES

(57) A medicament for the therapeutic treatment of diabetes which comprises as an active ingredient a compound represented by the general formula (I):



wherein R¹ represents a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aryl group, or a substituted or unsubstituted aromatic heterocyclic group, R² represents a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aralkyl group, a substituted or unsubstituted aryl group, or a substituted or unsubstituted aromatic heterocyclic group, R³ represents a hydrogen atom, a lower alkyl group or a substituted or unsubstituted aralkyl group, X¹ and X² independently represent a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aralkyl group, or a substituted or unsubstituted aryl group, and the symbol "n" represents an integer of from 0 to 3, or a physiologically acceptable salt thereof.

Description

Technical Field

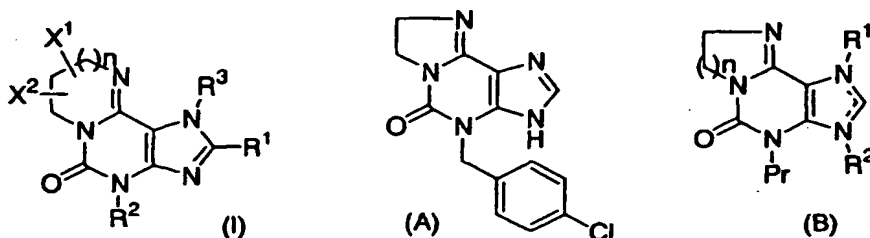
[0001] The present invention relates to an enhancer of insulin secretion and a medicament for the treatment of diabetes, which comprise a condensed purine derivative as an active ingredient.

Background Art

[0002] Diabetes is a metabolic abnormality mainly of glycometabolism, resulting from insufficient insulin secretion, decreased sensitivity of target cells of insulin and so forth, and principally characterized by noticeable hyperglycemia. If the hyperglycemia continues for a long period of time, serious complications arise in various organs and nerves such as retinopathy, nephropathy and neuropathy, which are caused mainly by vascular lesion. Therefore, for the treatment of diabetes, it is extremely important to control and maintain blood glucose level at a normal level, and methods for that purpose have been studied since old days.

[0003] For a type of diabetes where onset is gradual and insulin therapy is not necessarily required for life support (non-insulin dependent diabetes: NIDDM), blood glucose level can be controlled by combination of exercise therapy and drug therapy. As the drugs, insulin secretion enhancers, one of orally available hypoglycemic agents, have widely been used clinically. However, since currently available insulin secretion enhancers all promote insulin secretion non-dependently on glucose level, they cause problems of severe hypoglycemia or insufficient control of blood glucose if doses are not appropriate, and are not fully satisfactory drugs. If a hypoglycemic agent can be provided that is capable of enhancing insulin secretion dependently on a blood glucose level, the agent is expected to be extremely useful for blood glucose control of patients suffering from diabetes because the risk of hypoglycemia due to an excess dosage can be avoided.

[0004] As purine derivatives, the compounds of the following formula (I) having diuretic action, antiasthmatic action, anti-dementia action, bronchodilatation action, antiallergic action, antiulcer action and so forth are known (Japanese Patent Unexamined Publication (Kokai) No. 3-204880/1991; Journal of Medicinal Chemistry, vol. 35, p.3578, 1992; Journal of Medicinal Chemistry, vol. 36, p.2508, 1993; International Publication WO98/15555). Journal of Medicinal Chemistry, vol. 23, p.1188, 1980 discloses that the compounds represented by the formula (A) have bronchodilatation action. Furthermore, Journal of Medicinal Chemistry, vol. 40, p.3248, 1997 discloses that the compounds represented by the formula (B) have inhibitory action against type IV phosphodiesterase (bronchodilatation action).

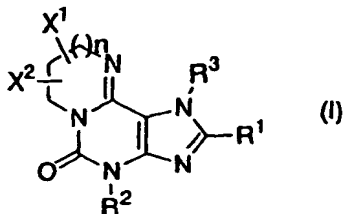


Disclosure of the Invention

[0005] An object of the present invention is to provide a medicament useful for the therapeutic treatment of diabetes and the prophylactic and/or therapeutic treatment of complications of diabetes. More specifically, the object of the present invention is to provide a medicament having action of enhancing insulin secretion in a blood glucose level-dependent manner.

[0006] The inventors of the present invention conducted various studies to achieve the foregoing object. As a result, we found that the compounds represented by the following general formula (I) have insulin secretion enhancing action, and that the compounds are useful as an active ingredient of a medicament for the treatment of diabetes. The present invention was achieved on the basis of these findings.

[0007] The present invention thus provides medicaments for the therapeutic treatment of diabetes which comprise compound represented by the general formula (I) or a physiologically acceptable salt thereof as an active ingredient:



wherein R¹ represents a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aryl group, or a substituted or unsubstituted aromatic heterocyclic group; R² represents a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aralkyl group, a substituted or unsubstituted aryl group, or a substituted or unsubstituted aromatic heterocyclic group; R³ represents a hydrogen atom, a lower alkyl group, or a substituted or unsubstituted aralkyl group; X¹ and X² independently represent a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aralkyl group, or a substituted or unsubstituted aryl group; and n represents an integer of from 0 to 3. The invention also provides medicaments for the prophylactic and/or therapeutic treatment of a complication of diabetes which comprise the aforementioned compound or a physiologically acceptable salt thereof as an active ingredient.

[0008] According to other aspects of the present invention, there are provided hypoglycemic agents comprising a compound represented by the aforementioned formula (I) or a physiologically acceptable salt thereof as an active ingredient, and insulin secretion enhancers comprising a compound represented by the aforementioned formula (I) or a physiologically acceptable salt thereof as an active ingredient. These medicaments are preferably provided in the form of a pharmaceutical composition which comprises a compound represented by the aforementioned formula (I) or a physiologically acceptable salt thereof and one or more of pharmaceutical additives.

[0009] According to other aspects of the present invention, there are provided use of a compound represented by the aforementioned formula (I) or a physiologically acceptable salt thereof for the manufacture of the aforementioned medicaments; a method for the therapeutic treatment of diabetes which comprises the step of administering a therapeutically effective amount of a compound represented by the aforementioned formula (I) or a physiologically acceptable salt thereof to a mammal including human; a method for enhancing insulin secretion which comprises the step of administering a therapeutically effective amount of a compound represented by the aforementioned formula (I) or a physiologically acceptable salt thereof to a mammal including human; and a method for decreasing blood glucose level which comprises the step of administering a therapeutically effective amount of a compound represented by the aforementioned formula (I) or a physiologically acceptable salt thereof to a mammal including human.

[0010] The compounds represented by the general formula (I) will be hereinafter referred to as Compound (I).

[0011] Meanings of terms used herein are as follows. As the lower alkyl group or a lower alkyl moiety of a functional group containing the lower alkyl moiety (for example, an aralkyl group), an alkyl group having about 1 to 10 carbon atoms can be used, and the term "alkyl group" includes linear, branched and cyclic alkyl groups as well as those consisting of a combination thereof. A cyclic alkyl group or a cyclic alkyl moiety of an alkyl group may have one or more rings, and may have, for example, a bicyclic ring or tricyclic ring system.

[0012] Examples of the linear or branched lower alkyl group include a methyl group, an ethyl group, a n-propyl group, an isopropyl group, a n-butyl group, an isobutyl group, a sec-butyl group, a tert-butyl group, a n-pentyl group, a neopentyl group, a n-hexyl group, a n-heptyl group, a n-octyl group, a n-nonyl group and so forth. Examples of the cyclic lower alkyl group include a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclooctyl group, a noradamantyl group, an adamantyl group and so forth.

[0013] Preferred examples of the lower alkyl group represented by R¹ include a linear or branched lower alkyl group (for example, a tert-butyl group, etc.), a monocyclic to tricyclic lower cycloalkyl group (for example, a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, a 3-noradamantyl group, etc.) and a lower alkyl group substituted with a lower cycloalkyl group (for example, a cyclopropylmethyl group, a cyclopentylmethyl group, etc.). Among them, a 3-noradamantyl group and a cyclopentyl group are more preferred. As the lower alkyl group represented by R², a linear or branched lower alkyl group (for example, a methyl group, an ethyl group, a n-propyl group, etc.), a monocyclic lower cycloalkyl group or a lower alkyl group substituted with a lower cycloalkyl group (for example, a cyclopropylmethyl group etc.) is preferred. Among them, a n-propyl group is more preferred. As the lower alkyl group represented by R³, a linear or branched lower alkyl group (for example, a methyl group, an ethyl group, etc.) is preferred, and a methyl group is more preferred. As the lower alkyl group represented by X¹ or X², a linear or branched lower alkyl group (for example, an ethyl group, an isopropyl group, a n-butyl group, an isobutyl group, a tert-butyl group, etc.), a monocyclic lower cycloalkyl group, or a lower alkyl group substituted with a lower cycloalkyl group (for example, a cyclohexylmethyl group etc.) is preferred. Among them, an ethyl group, an isopropyl group, an isobutyl group, and a tert-butyl group are more preferred.

[0014] As the aryl group or an aryl moiety of a functional group containing the aryl moiety (for example, an aralkyl group), an aromatic group consisting of a single ring or two or more condensed rings can be used. More specifically, an aryl group having 6 to 14 ring-membered carbon atoms is preferred. For example, a phenyl group, a

naphthyl group, an indenyl group, an anthranyl group or the like may preferably be used. As the aralkyl group, for example, an aralkyl group having 7 to 15 carbon atoms can be used. More specifically, a benzyl group, a phenethyl group, a diphenylmethyl group, a naphthylmethyl group or the like may be used.

[0015] As the aromatic heterocyclic group, an aromatic heterocyclic group consisting of a single ring or two or more condensed rings can be used. The kind and number of hetero atom(s) contained in the aromatic heterocyclic group are not particularly limited. For example, one or more hetero atoms selected from the group consisting of a nitrogen atom, a sulfur atom, and an oxygen atom may be contained. More specifically, an aromatic heterocyclic group containing about 6 to 14 ring-membered carbon atoms is preferred. For example, a furyl group, a thienyl group, a pyrrolyl group, an imidazolyl group, a pyrazolyl group, a triazolyl group, a tetrazolyl group, an oxazolyl group, a thiazolyl group, a pyridyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a triazinyl group, an indolyl group, an indazolyl group, a benzimidazolyl group, a benzoxazolyl group, a benzothiazolyl group, a quinolyl group, an isoquinolyl group, a phthalazinyl group, a naphthylidinyl group, a quinoxalinyl group, a quinazolinyl group, a cinnolinyl group, a purinyl group or the like may preferably be used.

[0016] When the aryl ring (including the aryl ring of an aralkyl group) or the aromatic heterocyclic ring has substituent(s), the kind, number and substitution position of the substituent(s) are not particularly limited. When two or more substituents exist, they may be the same or different. For example, the rings may have about 1 to 3 substituents. Examples of the substituents on the aryl ring or the aromatic heterocyclic ring include a substituted or unsubstituted lower alkyl group, a substituted or unsubstituted lower alkenyl group, a substituted or unsubstituted lower alkynyl group, a substituted or unsubstituted aralkyl group, a substituted or unsubstituted aryl group, a hydroxyl group, a substituted or unsubstituted lower alkoxy group, a substituted or unsubstituted aralkyloxy group, a substituted or unsubstituted aryloxy group, a substituted or unsubstituted aroyl group, a lower alkoxy carbonyl group, a substituted or unsubstituted lower alkylthio group, a substituted or unsubstituted lower alkylsulfonyl group, a carboxyl group, a substituted or unsubstituted carbamoyl group, a substituted or unsubstituted lower alkanoyl group, a halogen atom (the "halogen atom" used in the present specification may be any of a fluorine atom, a chlorine atom, a bromine atom, and an iodine atom), a nitro group, an amino group, a mono- or di(lower alkyl)-substituted amino group, a cyano group, a methylenedioxy group, an ethylenedioxy group and the like.

[0017] In the aforementioned functional groups, the above-explained groups may be used as the lower alkyl or a lower alkyl moiety of the functional groups containing a lower alkyl moiety (for example, the aralkyl group, the lower alkoxy group, the aralkyloxy group, the lower alkoxy carbonyl group, the lower alkylthio group, the lower alkylsulfonyl group, the lower alkanoyl group, the mono- or di(lower alkyl)-substituted amino group and the like). When the lower alkyl group or the lower alkyl moiety has substituent(s), the kind, number, and substitution position of the substituent(s) are not particularly limited. When two or more substituents exist, they may be the same or different. For example, the lower alkyl group or the lower alkyl moiety may have about 1 to 3 substituents. Examples of such substituents include a hydroxyl group, a halogen atom, a carboxyl group, a sulfo group, a phosphono group, ester groups derived from these acidic groups (a lower alkyl ester, an aralkyl ester, an aryl ester, etc.) and the like. More specific examples of the substituted alkyl group include a hydroxyethyl group, a trifluoromethyl group and the like. The two lower alkyl groups in the di(lower alkyl)-substituted amino group may be the same or different.

[0018] In the aforementioned functional groups, the aralkyl group or an aralkyl moiety of the aralkyloxy group has the same meaning as the aforementioned aralkyl group, and the aryl moiety of the aryl group, the aryloxy group, or the aroyl group has the same meaning as the aforementioned aryl group. As the lower alkenyl group, a linear or branched alkenyl group having 2 to 6 carbon atoms can be used. For example, a vinyl group, an allyl group, a 1-propenyl group, a methacryl group, a butenyl group, a crotyl group, a pentenyl group, a hexenyl group or the like may be used. As the lower alkynyl group, a linear or branched alkynyl group having 2 to 6 carbon atoms can be used. For example, an ethynyl group, a propynyl group, a butynyl group, a pentynyl group, a hexynyl group or the like may be used. The number of unsaturated bond in the alkenyl group or the alkynyl group is not particularly limited, and the number may preferably be 1.

[0019] When the aforementioned lower alkenyl group, lower alkynyl group, aralkyl group, aryl group, aralkyloxy group, aryloxy group, or aroyl group has substituent(s), the kind, number, and substitution position of the substituent(s) are not particularly limited. When two or more substituents exist, they may be the same or different. For example, the lower alkenyl group, lower alkynyl group, aralkyl group, aryl group, aralkyloxy group, aryloxy group, or aroyl group may have about 1 to 3 substituents. As the substituents, those exemplified as the substituents of the lower alkyl group may be used.

[0020] As the aryl group represented by R^1 , a phenyl group, a 4-bromophenyl group and the like are preferred. As the aromatic heterocyclic group represented by R^1 , a furyl group, a thienyl group and the like are preferred. As the aryl group represented by R^2 , a phenyl group is preferred. As the aralkyl group represented by R^2 , a benzyl group is preferred. As the aralkyl group represented by R^3 , a benzyl group is preferred. As the aralkyl group represented by X^1 or X^2 , a phenyl group is preferred. Examples of the aralkyl group represented by X^1 or X^2 include a benzyl group, a halogenated benzyl group (a 4-fluorobenzyl group, a 3-fluorobenzyl group, a 2-fluorobenzyl group, a 2,3-difluorobenzyl group, a 2,5-difluorobenzyl group, a 4-chlorobenzyl group, a 3-chlorobenzyl group, a 2,5-dichlorobenzyl group, a 4-bromobenzyl group, a 3-bromobenzyl group, an 3-iodobenzyl group, etc.), a 4-tert-butylbenzyl group, a 2-phenylbenzyl group, a 3-trifluoromethylbenzyl group, a 3-trifluoromethoxybenzyl group, a 3,5-bis(trifluoromethyl)benzyl group, a 4-hydroxybenzyl group, a 4-methoxybenzyl group, a 4-nitrobenzyl group, a 3-cyanobenzyl group, a 3-carboxybenzyl group, a 3-phenoxybenzyl group, a 3-(2-fluorophenoxy)benzyl group, a 4-benzyloxybenzyl group, a diphenylmethyl group, a phenethyl group, an α , α -dimethylbenzyl group and the like. When

X¹ is a group other than a hydrogen atom, X² is preferably a hydrogen atom. A combination of a lower alkyl group as X¹ (for example, a methyl group) and an aralkyl group as X² (for example, a benzyl group) is also preferred.

[0021] In the aforementioned general formula (I), the substitution positions of X¹ and X² are not particularly limited, and they may substitute at any positions on the ring. When X¹ or X² is a substituent other than a hydrogen atom, the absolute configuration of the carbon atom to which it binds may be in either a S- or R-configuration. The symbol "n" may preferably be 0.

[0022] As the active ingredient of the medicament of the present invention, physiologically acceptable salts of Compound (I) may be used. Such physiologically acceptable salts include acid addition salts such as inorganic acid salts and organic acid salts; base addition salts such as metal salts, ammonium salts and organic amine addition salts; amino acid addition salts and the like. Examples of the physiologically acceptable acid addition salts include inorganic acid salts such as hydrochlorides, sulphates and phosphates, and organic acid salts such as acetates, maleates, fumarates, tartrates, and citrates. Examples of the physiologically acceptable metal salts include alkali metal salts such as sodium salts and potassium salts, alkaline earth metal salts such as magnesium salts and calcium salts, as well as aluminum salts, zinc salts and the like. Examples of the physiologically acceptable organic amine addition salts include addition salts of organic amines such as morpholine, piperidine and the like. Examples of the physiologically acceptable amino acid addition salts include addition salts of lysine, glycine, phenylalanine and the like.

[0023] As the active ingredient of the medicament of the present invention, the aforementioned compounds in free forms or physiologically acceptable salts thereof, as well as any hydrates and solvates thereof may be used. The kind of solvents forming the solvates is not particularly limited so long as they are physiologically acceptable. For example, ethanol, acetone and the like may be used. The compounds represented by the formula (I) may have one or more asymmetric carbons. As the active ingredient of the medicament of the present invention, any substances falling within the scope of the general formula (I) may be used, including optical isomers or diastereoisomers in pure forms, any mixtures of isomers, racemates and the like. When the compounds represented by the general formula (I) contain one or more double bonds, each configuration thereof may be in either a Z- or E-configuration.

[0024] The methods for the preparation of typical compounds falling within the compounds represented by the formula (I) are specifically disclosed in International Publication WO98/15555, and are also explained in Japanese Patent Unexamined Publication No. 3-204880/1991; Journal of Medicinal Chemistry, vol. 35, p.3578, 1992; Journal of Medicinal Chemistry, vol. 36, p.2508, 1993; Journal of Heterocyclic Chemistry, vol. 30, p.241, 1993 and the like. Furthermore, the methods for preparing preferred compounds shown below will be specifically explained in detail in the examples of the present specification. Therefore, those skilled in the art can prepare any compounds falling within the scope of the general formula (I) according to the processes disclosed in the aforementioned references or the manufacturing processes specifically described in the present specification, or by suitably changing reagents and starting materials, and further by appropriately modifying or altering these processes, if necessary.

[0025] The intermediate compounds and target compounds obtained in these manufacturing processes can be isolated and purified by methods for purification conventionally used in the field of synthetic organic chemistry, for example, neutralization, filtration, extraction, washing, drying, concentration, recrystallization, various kinds of chromatography and the like. The intermediate compounds can also be used in subsequent reactions without particular purification. When salts of the compounds represented by the general formula (I) are prepared, the compounds in a free form may be dissolved or suspended in a suitable solvent, and a suitable acid or base is added to the mixture to form a salt, and then the resulting salts may be isolated and purified as required. It is also possible to convert a target substance obtained in a form of salt into a free form, and then converted into a desired salt.

[0026] Preferred examples of the active ingredient of the medicament of the present invention include:

(1) those wherein R¹ represents a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aryl group, or a substituted or unsubstituted aromatic heterocyclic group, R² represents a lower alkyl group, a substituted or unsubstituted aralkyl group or a substituted or unsubstituted aryl group, R³ represents a hydrogen atom, a lower alkyl group, or a substituted or unsubstituted aralkyl group, X¹ and X² independently represent a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aralkyl group or a substituted or unsubstituted aryl group, and the symbol "n" represents 0, or physiologically acceptable salts thereof; more preferred examples include:

(2) those wherein R¹ represents a lower alkyl group, a substituted or unsubstituted aryl group, or a substituted or unsubstituted aromatic heterocyclic group, R² represents a lower alkyl group or a substituted or unsubstituted aryl group, R³ represents a hydrogen atom or a substituted or unsubstituted aralkyl group, X¹ represents a hydrogen atom, a lower alkyl group, or a substituted or unsubstituted aralkyl group, X² represents a hydrogen atom, and the symbol "n" represents 0, or physiologically acceptable salts thereof; and further preferred examples include

(3) those wherein R¹ represents a n-propyl group, a n-butyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, a cyclopropylmethyl group, a cyclopentylmethyl group, an isopropyl group, or a tert-butyl group, R² represents a n-propyl group or an ethyl group, R³ represents a hydrogen atom, X¹ represents a lower alkyl

group or a substituted or unsubstituted aralkyl group, X² represents a hydrogen atom, and the symbol "n" represents 0, or physiologically acceptable salts thereof.

[0027] More specifically, the followings can be exemplified as preferred compounds or physiologically acceptable salts thereof suitable as the active ingredient of the medicament of the present invention. Among them, Compound 16, Compounds 18-20, Compound 23, Compound 31, Compound 35, Compound 37, Compound 43, and Compound 55 are particularly preferred. The compounds in free forms or other physiologically acceptable salts of these salts are also preferred. References disclosing each corresponding compound are indicated in the parentheses after the names of compounds. As for Compound 6, a racemate thereof is disclosed in Japanese Patent Unexamined Publication No. 3-204880/1991. It should be understood that the active ingredients of the medicaments of the present invention are not limited to the compounds or physiologically acceptable salts thereof listed below. The compounds specifically disclosed in Japanese Patent Unexamined Publication No. 3-204880/1991 or International Publication WO98/15555 or physiologically acceptable salts thereof, as well as those listed below, are preferred as the active ingredient of the medicament of the present invention.

Compound 1: (R)-8-ethyl-7,8-dihydro-2-(3-noradamantyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one L-tartrate (Journal of Medicinal Chemistry, vol. 36, p.2508, 1993)

Compound 2: (S)-8-ethyl-7,8-dihydro-2-(3-noradamantyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one D-tartrate (Journal of Medicinal Chemistry, vol. 36, p.2508, 1993)

Compound 3: (R)-7,8-dihydro-8-isopropyl-2-(3-noradamantyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 4: (S)-7,8-dihydro-8-isopropyl-2-(3-noradamantyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 5: 7,8-dihydro-2-(3-noradamantyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 6: (R)-2-cyclopentyl-7,8-dihydro-8-methyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 7: (R)-2-cyclopentyl-8-ethyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one L-tartrate (Japanese Patent Unexamined Publication No. 3-204880/1991)

Compound 8: (S)-2-cyclopentyl-8-ethyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one D-tartrate (Japanese Patent Unexamined Publication No. 3-204880/1991)

Compound 9: (R)-2-cyclopentyl-7,8-dihydro-8-isopropyl-4-propyl-1H-imidazo [2,1-i]purin-5(4H)-one hydrochloride (Japanese Patent Unexamined Publication 3-204880/1991)

Compound 10: (S)-2-cyclopentyl-7,8-dihydro-8-isopropyl-4-propyl-1H-imidazo [2,1-i]purin-5(4H)-one hydrochloride (Japanese Patent Unexamined Publication No. 3-204880/1991)

Compound 11: (R)-2-cyclopentyl-7,8-dihydro-8-isobutyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one L-tartrate

Compound 12: (S)-8-tert-butyl-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 13: 8-butyl-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 14: (S)-8-cyclohexylmethyl-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 15: (R)-2-cyclopentyl-7,8-dihydro-8-phenyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride (Japanese Patent Unexamined Publication No. 3-204880/1991)

Compound 16: (R)-8-benzyl-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride (Japanese Patent Unexamined Publication No. 3-204880/1991)

Compound 17: (S)-8-benzyl-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride (Japanese Patent Unexamined Publication No. 3-204880/1991)

Compound 18: 2-cyclopentyl-8-(4-fluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 19: 2-cyclopentyl-8-(3-fluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

hydrochloride

Compound 20: 8-(4-chlorobenzyl)-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 21: 8-(3-chlorobenzyl)-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 22: 2-cyclopentyl-8-(2,6-dichlorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 23: 8-(4-bromobenzyl)-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 24: 8-(3-bromobenzyl)-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 25: 2-cyclopentyl-7,8-dihydro-8-(4-methoxybenzyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 26: 2-cyclopentyl-7,8-dihydro-8-(4-nitrobenzyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 27: (R)-8(4-benzyloxybenzyl)-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 28: (S)-8-(4-benzyloxybenzyl)-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 29: (R)-8-benzyl-2-cyclopentyl-7,8-dihydro-1-methyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 30: 2-cyclopentyl-7,8-dihydro-8-diphenylmethyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 31: 2-cyclopentyl-7,8-dihydro-8-phenethyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 32: (R)-4,8-dibenzyl-2-cyclopentyl-7,8-dihydro-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 33: (R)-8-benzyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 34: (R)-1,8-dibenzyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 35: (R)-8-benzyl-7,8-dihydro-2,4-dipropyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 36: (R)-8-benzyl-2-cyclopropyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 37: (R)-8-benzyl-2-cyclobutyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 38: (R)-8-benzyl-2-cyclohexyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 39: (R)-8-benzyl-7,8-dihydro-2-(3-noradamantyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 40: (R)-8-benzyl-2-cyclopentylmethyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 41: (R)-8-benzyl-7,8-dihydro-2-isopropyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 42: (R)-8-benzyl-2-(3-furyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 43: (R)-8-benzyl-2-tert-butyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 44: (R)-8-benzyl-7,8-dihydro-4-propyl-2-(3-thienyl)-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 45: 8-(4-bromobenzyl)-2-cyclohexyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 46: 8-(4-chlorobenzyl)-7,8-dihydro-2,4-dipropyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

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Compound 47: 8-(4-chlorobenzyl)-2-cyclohexyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 48: 8-(4-chlorobenzyl)-2-cyclobutyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 49: 8-(4-fluorobenzyl)-7,8-dihydro-2,4-dipropyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 50: 2-cyclohexyl-8-(4-fluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 51: (R)-2-cyclobutyl-8-(4-fluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 52: 2-butyl-8-(4-fluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 53: 8-(4-fluorobenzyl)-2-(2-furyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 54: (R)-8-(4-fluorobenzyl)-2-(3-furyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 55: (R)-2-tert-butyl-8-(4-fluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 56: 8-(3-fluorobenzyl)-7,8-dihydro-2,4-dipropyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 57: 2-cyclobutyl-8(3-fluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 58: 8-(3-fluorobenzyl)-7,8-dihydro-2-isobutyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 59: 2-(4-bromophenyl)-8-(3-fluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 60: 8-(2-fluorobenzyl)-7,8-dihydro-2-isobutyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 61: 2-cyclopentyl-7,8-dihydro-8-(3-methylbenzyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 62: 2-cyclopentyl-7,8-dihydro-8-(3-iodobenzyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 63: 2-cyclopentyl-8-(2,3-difluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 64: 2-cyclopentyl-8-(2,5-difluorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 65: 2-cyclopentyl-8-(2,5-dichlorobenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 66: (S)-8-tert-butyl-7,8-dihydro-2-(3-noradamantyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 67: (S)-8-tert-butyl-2-(3-furyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 68: (S)-8-tert-butyl-2-(3,5-dimethylisoxazol-4-yl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 69: (S)-7,8-dihydro-2,8-diisopropyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 70: 8-(3-cyanobenzyl)-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 71: (R)-8-benzyl-2-cyclopentyl-4-ethyl-7,8-dihydro-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 72: (R)-8-benzyl-2-cyclopentyl-4-cyclopropylmethyl-7,8-dihydro-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 73: (R)-8-benzyl-2-cyclopropylmethyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one hydrochloride

Compound 74: 8-(3-carboxybenzyl)-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

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Compound 75: 2-cyclopentyl-8-(α , α -dimethylbenzyl)-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 76: (R)-2-cyclopentyl-7,8-dihydro-8-(4-hydroxybenzyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 77: (R)-8-benzyl-7,8-dihydro-4-propyl-2-(3-pyridyl)-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 78: (R)-8-benzyl-7,8-dihydro-2-phenyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 79: (R)-8-benzyl-7,8-dihydro-4-propyl-2-(2,2,3,3-tetramethylcyclopropyl)-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 80: (R)-8-benzyl-7,8-dihydro-2-(3,4-methylenedioxyphenyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 81: (R)-8-benzyl-7,8-dihydro-4-propyl-2-(4-pyridyl)-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 82: (R)-8-benzyl-7,8-dihydro-4-propyl-2-(2-pyrazyl)-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 83: (R)-8-benzyl-7,8-dihydro-4-propyl-2-(2-thienyl)-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 84: (R)-8-benzyl-2-cyclopentyl-7,8-dihydro-4-methyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 85: (R)-8-benzyl-2-cyclopentyl-7,8-dihydro-4-phenyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 86: 8-(4-tert-butylbenzyl)-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 87: 2-cyclopentyl-7,8-dihydro-8-(2-phenylbenzyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 88: 2-cyclopentyl-7,8-dihydro-4-propyl-8-(3-trifluoromethylbenzyl)-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 89: 2-cyclopentyl-8-[3-(2-fluorophenoxy)benzyl]-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

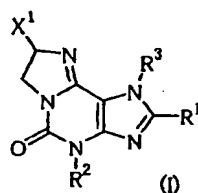
Compound 90: 2-cyclopentyl-7,8-dihydro-8-(3-phenoxybenzyl)-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 91: 2-cyclopentyl-7,8-dihydro-4-propyl-8-(3-trifluoromethoxybenzyl)-1H-imidazo[2,1-i]purin-5(4H)-one

Compound 92: 8-[3,5-bis(trifluoromethyl)benzyl]-2-cyclopentyl-7,8-dihydro-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

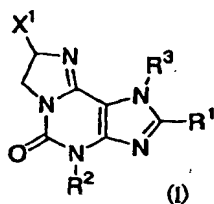
Compound 93: 8-benzyl-2-cyclopentyl-7,8-dihydro-8-methyl-4-propyl-1H-imidazo[2,1-i]purin-5(4H)-one

Table 1 (I)



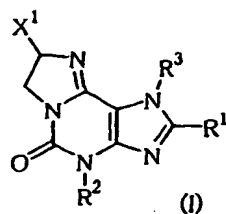
Compound No.	R ¹	R ²	R ³	X ¹
1		(CH ₂) ₂ CH ₃	H	H ₃ C-
2		(CH ₂) ₂ CH ₃	H	H ₃ C-
3		(CH ₂) ₂ CH ₃	H	H ₃ C--CH ₃
4		(CH ₂) ₂ CH ₃	H	H ₃ C--CH ₃
5		(CH ₂) ₂ CH ₃	H	H
6		(CH ₂) ₂ CH ₃	H	H ₃ C-
7		(CH ₂) ₂ CH ₃	H	H ₃ C-
8		(CH ₂) ₂ CH ₃	H	H ₃ C-
9		(CH ₂) ₂ CH ₃	H	H ₃ C--CH ₃
10		(CH ₂) ₂ CH ₃	H	H ₃ C--CH ₃
11		(CH ₂) ₂ CH ₃	H	H ₃ C--CH ₃ -CH ₃
12		(CH ₂) ₂ CH ₃	H	H ₃ C--CH ₃ -CH ₃ -
13		(CH ₂) ₂ CH ₃	H	H ₃ C-
14		(CH ₂) ₂ CH ₃	H	

Table 1 (2)



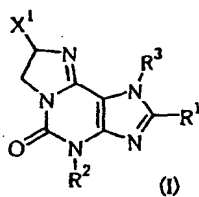
Compound No.	R ¹	R ²	R ³	X ¹
15		(CH ₂) ₂ CH ₃	H	
16		(CH ₂) ₂ CH ₃	H	
17		(CH ₂) ₂ CH ₃	H	
18		(CH ₂) ₂ CH ₃	H	
19		(CH ₂) ₂ CH ₃	H	
20		(CH ₂) ₂ CH ₃	H	
21		(CH ₂) ₂ CH ₃	H	
22		(CH ₂) ₂ CH ₃	H	
23		(CH ₂) ₂ CH ₃	H	
24		(CH ₂) ₂ CH ₃	H	
25		(CH ₂) ₂ CH ₃	H	
26		(CH ₂) ₂ CH ₃	H	

Table 1 (3)



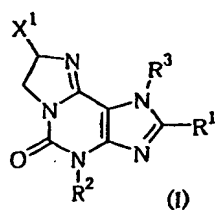
Compound No.	R ¹	R ²	R ³	X ¹
27		(CH ₂) ₂ CH ₃	H	
28		(CH ₂) ₂ CH ₃	H	
29		(CH ₂) ₂ CH ₃	CH ₃	
30		(CH ₂) ₂ CH ₃	H	
31		(CH ₂) ₂ CH ₃	H	
32				

Table 1 (4)



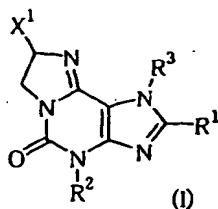
Compound No.	R^1	R^2	R^3	X^1
33	H	$(CH_2)_2CH_3$	H	
34	H	$(CH_2)_2CH_3$		
35		$(CH_2)_2CH_3$	H	
36		$(CH_2)_2CH_3$	H	
37		$(CH_2)_2CH_3$	H	
38		$(CH_2)_2CH_3$	H	
39		$(CH_2)_2CH_3$	H	
40		$(CH_2)_2CH_3$	H	
41		$(CH_2)_2CH_3$	H	
42		$(CH_2)_2CH_3$	H	
43		$(CH_2)_2CH_3$	H	
44		$(CH_2)_2CH_3$	H	
45		$(CH_2)_2CH_3$	H	

Table 1 (5)



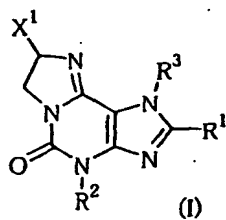
Compound No.	R^1	R^2	R^3	X^1
46		$(CH_2)_2CH_3$	H	
47		$(CH_2)_2CH_3$	H	
48		$(CH_2)_2CH_3$	H	
49		$(CH_2)_2CH_3$	H	
50		$(CH_2)_2CH_3$	H	
51		$(CH_2)_2CH_3$	H	
52		$(CH_2)_2CH_3$	H	
53		$(CH_2)_2CH_3$	H	
54		$(CH_2)_2CH_3$	H	
55		$(CH_2)_2CH_3$	H	
56		$(CH_2)_2CH_3$	H	
57		$(CH_2)_2CH_3$	H	

Table 1 (6)



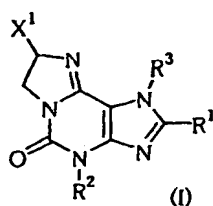
Compound No.	R ¹	R ²	R ³	X ¹
58		(CH ₂) ₂ CH ₃	H	
59		(CH ₂) ₂ CH ₃	H	
60		(CH ₂) ₂ CH ₃	H	
61		(CH ₂) ₂ CH ₃	H	
62		(CH ₂) ₂ CH ₃	H	
63		(CH ₂) ₂ CH ₃	H	
64		(CH ₂) ₂ CH ₃	H	
65		(CH ₂) ₂ CH ₃	H	

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Table 1 (7)



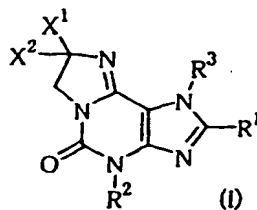
Compound No.	R ¹	R ²	R ³	X ¹
66		(CH ₂) ₂ CH ₃	H	
67		(CH ₂) ₂ CH ₃	H	
68		(CH ₂) ₂ CH ₃	H	
69		(CH ₂) ₂ CH ₃	H	
70		(CH ₂) ₂ CH ₃	H	
71		CH ₂ CH ₃	H	
72			H	
73		(CH ₂) ₂ CH ₃	H	
74		(CH ₂) ₂ CH ₃	H	

Table 1 (8)



Compound No.	R ¹	R ²	R ³	X ¹
75		(CH ₂) ₂ CH ₃	H	
76		(CH ₂) ₂ CH ₃	H	
77		(CH ₂) ₂ CH ₃	H	
78		(CH ₂) ₂ CH ₃	H	
79		(CH ₂) ₂ CH ₃	H	
80		(CH ₂) ₂ CH ₃	H	
81		(CH ₂) ₂ CH ₃	H	
82		(CH ₂) ₂ CH ₃	H	
83		(CH ₂) ₂ CH ₃	H	
84		CH ₃	H	
85		C ₆ H ₅	H	

Table 1 (9)



Compound No.	R ¹	R ²	R ³	X ¹	X ²
86		(CH ₂) ₂ CH ₃	H		H
87		(CH ₂) ₂ CH ₃	H		H
88		(CH ₂) ₂ CH ₃	H		H
89		(CH ₂) ₂ CH ₃	H		H
90		(CH ₂) ₂ CH ₃	H		H
91		(CH ₂) ₂ CH ₃	H		H
92		(CH ₂) ₂ CH ₃	H		H
93		(CH ₂) ₂ CH ₃	H	CH ₃	

[0028] The compounds represented by the general formula (I) or physiologically acceptable salts thereof have insulin secretion enhancing action in cultured β -cells and hypoglycemic action in rats, and accordingly, they are useful as an active ingredient of the medicament for the treatment of diabetes. Further, they are useful as an active ingredient of the medicament for the prophylactic and/or therapeutic treatment of various complications of diabetes, for example, retinopathy, nephropathy, neurosis and the like. As the active ingredient of the medicament of the present invention, one or more substances selected from the group consisting of the compounds represented by

the general formula (I) and physiologically acceptable salts thereof, and hydrates thereof and solvates thereof can be used. Although the aforementioned substances, per se, may be administered, it is generally desirable to provide the medicament in a form of a pharmaceutical composition comprising the aforementioned substance as the active ingredient and one or more pharmaceutical additives. The medicament of the present invention can be administered to human and other mammals.

[0029] The form of the pharmaceutical composition is not particularly limited, and an appropriate form most suitable for a purpose of the prophylactic or therapeutic treatment can be selected from forms of pharmaceutical preparations for oral or parenteral administration. Examples of pharmaceutical preparations suitable for oral administration include tablets, capsules, powders, granules, subitized granules, syrups, solutions, emulsions, suspensions, chewable formulations and the like. Examples of pharmaceutical preparations suitable for parenteral administration include injections (for subcutaneous injection, intramuscular injection, intravenous injection or the like), drip infusions, inhalants, sprays, suppositories, transdermal or transmucosal preparations in the forms of gel, ointment or the like, transdermal preparations in the forms of patch, tape or the like. However, the preparations are not limited to these examples.

[0030] Liquid preparations suitable for oral administration such as solutions, emulsions and syrups can be prepared by using water, saccharides such as sucrose, sorbitol and fructose, glycols such as polyethylene glycol and propylene glycol, oils such as sesame oil, olive oil and soybean oil, preservatives such as p-hydroxybenzoic acid esters, flavors such as strawberry flavor and peppermint or the like. For the preparation of solid preparations such as capsules, tablets, powders and granules, excipients such as lactose, glucose, sucrose and mannitol, disintegrating agents such as starch and sodium alginate, lubricants such as magnesium stearate and talc, binders such as polyvinyl alcohol, hydroxypropylcellulose and gelatin, surface active agents such as fatty acid esters, plasticizers such as glycerin or the like may be used.

[0031] Pharmaceutical preparations for injection or drip infusion, which are suitable for parenteral administration, contain the aforementioned substance as the active ingredient preferably in a sterilized aqueous medium isotonic with blood of a recipient in a dissolved or suspended state. For example, as for injections, a solution can be prepared by using an aqueous medium consisting of saline, a glucose solution, or a mixture of saline and a glucose solution. Preparations for enteral administration can be prepared, for example, by using a carrier such as cacao butter or hard fat, and are provided as suppositories. Further, for the preparation of sprays, a carrier may be used which can disperse the aforementioned substance as the active ingredient as fine particles, and facilitate adsorption of the active ingredient without stimulating mucosae of oral cavity and respiratory tract of recipients. Specifically, examples of the carrier include lactose and glycerol. Depending on the properties of the substance as the active ingredient and the carrier, pharmaceutical preparations in forms of aerosol, dry powder and the like can be prepared. To these pharmaceutical preparations for parenteral administration, one or more auxiliary ingredients selected from glycols, oils, flavors, preservatives, excipients, disintegrating agents, lubricants, binders, surface active agents, plasticizers and the like may also be added.

[0032] Dose and frequency of administration of the medicament of the present invention may preferably be increased or decreased depending on various factors such as type and severity of diseases, dosage form, conditions of patients such as age and body weight, and presence or absence of complications. In general, the medicament may preferably be administered in an amount of 1 to 1000 mg/kg per day for an adult dividedly as three or four times of administrations.

[0033] According to another aspect of the present invention, there are provided the compounds of the general formula (I):

(1) wherein R^1 represents a 3-noradamantyl group, R^2 represents a n-propyl group, R^3 represents a hydrogen atom, X^1 represents a hydrogen atom, a tert-butyl group, an isopropyl group, or a benzyl group, X^2 represents a hydrogen atom, and the symbol "n" represents 0;

(2) wherein R^1 represents a cyclopentyl group, R^2 represents a n-propyl group, R^3 represents a hydrogen atom, X^1 represents a n-butyl group, an isobutyl group, a tert-butyl group, a cyclohexylmethyl group, an aralkyl group whose alkyl moiety is substituted or an unsubstituted aralkyl group (except for a benzyl group), or an aralkyl group (preferably a benzyl group) having one or more substituents (preferably one or two, more preferably one) selected from the group consisting of a halogen atom, a substituted or unsubstituted lower alkyl group, a lower alkoxy group (preferably a methoxy group), an aralkyloxy group (preferably a benzyloxy group), a carboxyl group, a cyano group, a hydroxyl group, and a nitro group on the ring, X^2 represents a hydrogen atom, and the symbol "n" represents 0;

(3) wherein R^1 represents a cyclopentyl group, R^2 represents a n-propyl group, R^3 represents a lower alkyl group (preferably a methyl group), X^1 represents a substituted or unsubstituted aralkyl group (preferably a benzyl group), X^2 represents a hydrogen atom, and the symbol "n" represents 0;

(4) wherein R^1 represents a cyclopentyl group, R^2 represents a substituted or unsubstituted aralkyl group (preferably a benzyl group), a lower alkyl group (except for a n-propyl group; preferably a cyclopropylmethyl group or a linear lower alkyl group) or a substituted or unsubstituted aryl group (preferably a phenyl group),

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R³ represents a hydrogen atom, X¹ represents a substituted or unsubstituted aralkyl group (preferably a benzyl group), X² represents a hydrogen atom, and the symbol "n" represents 0;

(5) wherein R¹ represents a hydrogen atom, R² represents a n-propyl group, R³ represents a hydrogen atom, or a substituted or unsubstituted aralkyl group (preferably a benzyl group), X¹ represents a substituted or unsubstituted aralkyl group (preferably a benzyl group), X² represents a hydrogen atom, and the symbol "n" represents 0;

(6) wherein R¹ represents a lower alkyl group (except for a cyclopentyl group), a substituted or unsubstituted aromatic heterocyclic group, or a substituted or unsubstituted aryl group, R² represents a n-propyl group, R³ represents a hydrogen atom, X¹ represents a substituted or unsubstituted aralkyl group (preferably a benzyl group), X² represents a hydrogen atom, and the symbol "n" represents 0;

(7) wherein R¹ represents a linear or branched lower alkyl group (preferably an isopropyl group) or a substituted or unsubstituted aromatic heterocyclic group, R² represents a n-propyl group, R³ represents a hydrogen atom, X¹ represents a lower alkyl group (except for a methyl group; preferably an isopropyl group, or an isobutyl group, a t-butyl group), X² represents a hydrogen atom, and the symbol "n" represents 0; and

(8) wherein R¹ represents a cyclopentyl group, R² represents a n-propyl group, R³ represents a hydrogen atom, X¹ represents a methyl group, X² represents a substituted or unsubstituted aralkyl group, and the symbol "n" represents 0.

[0034] Specific examples of these compounds include, for example, Compounds 3-5, Compounds 11-14, and Compound 18 to Compound 93.

[0035] As preferred compounds, there are provided:

the aforementioned compounds wherein R¹ represents a cyclopentyl group, R² represents a n-propyl group, R³ represents a hydrogen atom, Xi represents an aralkyl group (preferably a benzyl group) having one or more (preferably one) halogen atoms on the ring or an unsubstituted aralkyl group (except for a benzyl group), X² represents a hydrogen atom, and the symbol "n" represents 0;

the aforementioned compounds wherein R¹ represents a cyclopentyl group, R² represents a linear or branched lower alkyl group (except for a n-propyl group), R³ represents a hydrogen atom, X¹ represents a substituted or unsubstituted aralkyl group (preferably a benzyl group), X² represents a hydrogen atom, and the symbol "n" represents 0; and

the aforementioned compounds wherein R¹ represents a lower alkyl group (except for a cyclopentyl group), R² represents a n-propyl group, R³ represents a hydrogen atom, X¹ represents a substituted or unsubstituted aralkyl group (preferably a benzyl group), X² represents a hydrogen atom, and the symbol "n" represents 0;

[0036] Particularly preferred compounds are:

the aforementioned compounds where R¹ represents a cyclopentyl group, R² represents an ethyl group, R³ represents a hydrogen atom, X¹ represents a substituted or unsubstituted aralkyl group (preferably a benzyl group), X² represents a hydrogen atom, and the symbol "n" represents 0; and

the aforementioned compounds wherein R¹ represents a tert-butyl group, R² represents a n-propyl group, R³ represents a hydrogen atom, X¹ represents a substituted or unsubstituted aralkyl group (preferably a benzyl group), X² represents a hydrogen atom, and the symbol "n" represents 0.

[0037] As salts of the aforementioned compounds, those explained above can be used. It should be understood that, besides the aforementioned compounds in free forms and physiologically acceptable salts thereof, any hydrates and solvates thereof fall within the scope of the present invention. Further, any isomers, mixtures of isomers, racemates and the like also fall within the scope of the present invention. These compounds and salts thereof are useful as, for example, the active ingredient of the medicament of the present invention. However, utilities of the

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compounds of the present invention and salts thereof are not limited to said particular utility.

Best Mode for Carrying out the Invention

Examples

[0038] The present invention will be explained more specifically with reference to the following examples. However, the scope of the present invention is not limited to the following examples.

Formulation Example 1: Tablet

[0039] Tablets consisting of the following composition were prepared in a conventional manner.

Formulation

[0040]

Compound 16	20 mg
Lactose	143.4 mg
Potato starch	30 mg
Hydroxypropylcellulose	6 mg
Magnesium stearate	0.6 mg
	200 mg

Formulation Example 2: Capsule

[0041] Capsules consisting of the following composition were prepared in a conventional manner.

Formulation

[0042]

Compound 16	20 mg
Avicell	99.5 mg
Magnesium stearate	0.5 mg
	120 mg

Formulation Example 3: Injection

[0043] Injections consisting of the following composition were prepared in a conventional manner.

Formulation

[0044]

Compound 16	2 mg
Purified soybean oil	200 mg
Purified yolk lecithin	24 mg
Glycerol for injection	50 mg
Distilled water for injection	1.72 ml
	2.00 ml

Formulation Example 4: Suppository for anus

[0045] Preparations for rectal administration consisting of the following composition were prepared in a conventional manner.

Formulation

[0046]

Compound 16	2.5 mg
Witepsol H15	678.8 mg
Witepsol E75	290.9 mg
Potassium primary phosphate	13.6 mg
Sodium secondary phosphate	14.2 mg
	1,000 mg

Example 1: Compound 3

[0047] To 6-methylthio-8-(3-noradamantyl)-3-propyl-7H-purin-2(3H)-one (Compound 3a, 3.06 g, 8.90 mmol) obtained by a known method [Journal of Medicinal Chemistry, vol. 36, p.2508, 1993] was added (R)-valinol (5.7 ml, a 4.66 M solution in dimethyl sulfoxide, 3.0 eq.) under an argon flow, and the mixture was heated at 150°C for 3.5 hours with stirring. The reaction solution was poured into a 10% aqueous solution of ammonium chloride, and the mixture was extracted with chloroform. The organic layer was washed with a saturated aqueous solution of sodium chloride and dried, and then the solvent was evaporated. The residue was purified by silica gel column chromatography (chloroform:methanol = 10:1) to obtain an adduct (2.97 g, 84 %). To the adduct (2.78 g, 6.97 mmol) was added thionyl chloride (30 ml), and the mixture was heated at 60°C for 3 hours with stirring. After the excessive thionyl chloride was evaporated, the reaction mixture was neutralized with sodium hydrogen carbonate and extracted with chloroform. The organic layer was washed with a saturated aqueous solution of sodium chloride and dried, and the solvent was evaporated. The residue was purified by silica gel column chromatography (chloroform:methanol = 50:1), and the solvent was evaporated. A methanolic solution of hydrochloric acid was added to the residue and the solvent was evaporated. The residue was crystallized from ethyl acetate/hexane to obtain Compound 3 (2.25 g, 77 %).

Yield: 65%

Melting point: 105-107°C (cyclohexane)

Optical rotation: $[\alpha]_D^{20} +36.9^\circ$ {c 1.00, CH₃OH, >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]}

¹H NMR (270 MHz, CDCl₃) δ : 0.98 (t, 3H, J=7.4Hz), 1.05 (d, 3H, J=6.6Hz), 1.14 (d, 3H, J=7.0Hz), 1.65-2.10 (m, 11H), 2.30 (m, 2H), 2.41 (m, 2H), 2.69 (t, 1H, J=6.0Hz), 4.01 (dd, 1H, J=11.0, 7.0Hz), 4.09-4.22 (m, 3H), 4.32 (dd, 1H, J=11.0, 10.0Hz)

IR (KBr): 1714, 1688, 1591 cm⁻¹

EI-MS: m/e 381 (M⁺)

Elemental analysis: for C ₂₂ H ₃₁ N ₅ O ₄ HCl \cdot 0.4H ₂ O			
Calculated (%):	C, 63.87;	H, 8.26;	N, 15.26
Found (%):	C, 63.92;	H, 8.19;	N, 15.20

Example 2: Compound 4

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[0048] Compound 4 was obtained from Compound 3a and (S)-valinol in the same manner as Example 1.

Yield: 57%

Melting point: 108-110°C (cyclohexane)

Optical rotation: $[\alpha]_D^{20}$ -30.8° (c 1.00, CH₃OH, >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

Elemental analysis: for C ₂₂ H ₃₁ N ₅ O ₄ HCl _{0.25} C ₆ H ₁₂			
Calculated (%):	C, 64.29;	H, 8.04;	N, 15.95
Found (%):	C, 63.72;	H, 8.30;	N, 15.43

Example 3: Compound 5

[0049] Compound 5 was obtained from Compound 3a and 2-aminoethanol in the same manner as Example 1.

Yield: 21%

Melting point: 274-276°C (ethyl acetate/toluene)

¹H NMR (270 MHz, CDCl₃) δ : 0.99 (t, 3H, J=7.4Hz), 1.67-2.08 (m, 12H), 2.20-2.28 (m, 2H), 2.40-2.43 (m, 2H), 2.71 (t, 1H, J=6.0Hz), 4.09-4.20 (m, 4H), 4.27-4.40 (m, 2H)

IR (KBr): 1719, 1669, 1590 cm⁻¹

EI-MS: m/e 339 (M⁺)

Elemental analysis: for C ₁₉ H ₂₅ N ₅ O ₄ HCl _{0.5} H ₂ O			
Calculated (%):	C, 59.29;	H, 7.07;	N, 18.19
Found (%):	C, 59.31;	H, 7.24;	N, 18.07

Example 4: Compound 6

[0050] Compound 6 was obtained from 8-cyclopentyl-6-methylthio-3-propyl-7H-purin-2(3H)-one (Compound 6a, Journal of Heterocyclic Chemistry, vol. 30, p.241, 1993) and (R)-alaninol in the same manner as Example 1.

Yield: 43%

Melting point: 218-219°C (ethyl acetate/cyclohexane)

Optical rotation: $[\alpha]_D^{20}$ +41.5° (c 1.00, CH₃OH, 96% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

Elemental analysis: for C ₁₆ H ₂₃ N ₅ O ₄ HCl _{0.2} H ₂ O			
Calculated (%):	C, 56.28;	H, 7.20;	N, 20.51
Found (%):	C, 56.14;	H, 7.45;	N, 20.33

Example 5: Compound 7

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[0051] Compound 7 was obtained from Compound 6a and (R)-2-amino-1-butanol in the same manner as Example 1 except that the compound was obtained as L-tartrate instead of hydrochloride.

Yield: 18%

Melting point: 209°C (isopropanol)

Optical rotation: $[\alpha]_D^{20} +4.2^\circ$ (c 1.00, dioxane, 95% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

Elemental analysis: for $C_{16}H_{23}N_5O_4C_4H_6O_6$			
Calculated (%):	C, 54.18;	H, 6.71;	N, 15.04
Found (%):	C, 54.47;	H, 6.60;	N, 14.68

Example 6: Compound 8

[0052] Compound 8 was obtained from Compound 6a and (S)-2-amino-1-butanol in the same manner as Example 1 except that the compound was obtained as D-tartrate instead of hydrochloride.

Yield: 9%

Melting point: 210-212°C (isopropanol)

Optical rotation: $[\alpha]_D^{20} -5.3^\circ$ (c 1.00, dioxane, >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

Elemental analysis: for $C_{16}H_{23}N_5O_4C_4H_6O_6$			
Calculated (%):	C, 54.18;	H, 6.71;	N, 15.04
Found (%):	C, 53.74;	H, 6.60;	N, 14.88

Example 7: Compound 9

[0053] Compound 9 was obtained from Compound 6a and (R)-valinol in the same manner as Example 1.

Yield: 26%

Melting point: 116-118°C (toluene/cyclohexane)

Optical rotation: $[\alpha]_D^{20} +51.1^\circ$ (c 1.00, CH_3OH , >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

1H NMR (270 MHz, $CDCl_3$) δ : 0.90-1.15 (m, 9H), 1.60-2.00 (m, 6H), 2.10-2.25 (m, 4H), 3.20-3.40 (m, 1H), 4.00-4.40 (m, 6H), 11.32 (brs, 1H)

IR (KBr): 1719, 1717, 1681, 1593 cm^{-1}

EI-MS: m/e 329 (M^+)

Elemental analysis: for $C_{18}H_{27}N_5O_4 \cdot 0.5HCl \cdot 0.5H_2O$			
Calculated (%):	C, 60.70;	H, 7.92;	N, 19.66
Found (%):	C, 60.61;	H, 8.24;	N, 19.43

Example 8: Compound 10

[0054] Compound 10 was obtained from Compound 6a and (S)-valinol in the same manner as Example 1.

Yield: 61%

Melting point: 109-112°C (toluene/cyclohexane)

Optical rotation: $[\alpha]_D^{20}$ -53.9° (c 1.00, CH₃OH, >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

Elemental analysis: for C ₁₈ H ₂₇ N ₅ O ₆ ·0.5HCl·0.5H ₂ O			
Calculated (%):	C, 59.12;	H, 8.13;	N, 19.15
Found (%):	C, 59.11;	H, 8.29;	N, 18.81

Example 9: Compound 11

[0055] Compound 11 was obtained from Compound 6a and (R)-leucinol in the same manner as Example 5.

Yield: 65%

Melting point: 221-223°C (methanol/ether)

Optical purity: >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]

¹H NMR (270 MHz, DMSO-d₆) δ : 0.87 (t, 3H, J=7.6Hz), 0.91 (d, 3H, J=3.6Hz), 0.93 (d, 3H, J=4.3Hz), 1.42 (m, 1H), 1.44-1.84 (m, 10H), 1.87-1.97 (m, 2H), 3.12 (quin, 1H, J=8.1Hz), 3.56 (dd, 1H, J=10.9, 6.9Hz), 3.87 (dd, 2H, J=7.9, 6.9Hz), 4.12 (dd, 1H, J=10.9, 9.6Hz), 4.24 (m, 1H), 4.25 (s, 2H)

IR (CHCl₃): 1716, 1682, 1581 cm⁻¹

EI-MS: m/e 343 (M⁺)

Elemental analysis: for C ₁₉ H ₂₉ N ₅ O ₆ ·C ₄ H ₆ O ₆ ·0.4H ₂ O			
Calculated (%):	C, 55.17;	H, 7.21;	N, 13.99
Found (%):	C, 55.23;	H, 7.24;	N, 13.87

Example 10: Compound 12

[0056] Compound 12 was obtained from Compound 6a and (S)-tert-leucinol in the same manner as Example 1.

Yield: 47%

Melting point: 184-186°C (acetone/ether)

Optical rotation: $[\alpha]_D^{24}$ -33.3° (c 1.12, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

¹H NMR (270 MHz, CDCl₃) δ : 0.99 (t, 3H, J=7.4Hz), 1.06 (s, 9H), 1.65-1.95 (m, 8H), 2.10-2.22 (m, 2H), 3.28

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(quin, 1H, J=7.9Hz), 4.09 (m, 1H), 4.10 (t, 2H, J=7.6Hz), 4.23 (m, 2H)

IR (CHCl₃): 1717, 1687, 1587 cm⁻¹

ElúMS: m/e 343 (M⁺)

Elemental analysis: for C ₁₉ H ₂₉ N ₅ OúHClú0.5H ₂ O			
Calculated (%):	C, 58.67;	H, 8.03;	N, 18.01
Found (%):	C, 58.85;	H, 8.17;	N, 17.92

Example 11: Compound 13

[0057] Compound 13 was obtained from Compound 6a and 2-amino-1-hexanol in the same manner as Example 1.

Yield: 40%

Melting point: 146-148°C (acetone)

¹H NMR (270 MHz, CDCl₃) δ : 0.94 (t, 3H, J=7.1Hz), 0.98 (t, 3H, J=7.4Hz), 1.36-1.45 (m, 2H), 1.61-1.92 (m, 12H), 2.08-2.21 (m, 2H), 3.27 (quin, 1H, J=7.9Hz), 3.94 (m, 1H), 4.10 (t, 2H, J=7.4Hz), 4.34-4.41 (m, 2H)

IR (CHCl₃): 1732, 1716, 1699, 1687, 1506 cm⁻¹

EI-MS: m/e 343 (M⁺)

Elemental analysis: for C ₁₉ H ₂₉ N ₅ Oú1.2HCl			
Calculated (%):	C, 58.93;	H, 7.86;	N, 18.09
Found (%):	C, 59.22;	H, 7.96;	N, 18.13

Example 12: Compound 14

[0058] Compound 14 was obtained from Compound 6a and (S)-2-amino-3-cyclohexyl-1-propanol in the same manner as Example 1.

Yield: 36%

Melting point: 198-200°C (acetone/ether)

Optical rotation: [α]_D²⁴ -51.3° (c 0.60, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

¹H NMR (270 MHz, CDCl₃) δ : 0.98 (t, 3H, J=7.4Hz), 1.14-1.37 (m, 3H), 1.54-1.89 (m, 18H), 2.16 (m, 2H), 3.27 (quin, 1H, J=7.9Hz), 3.91 (dd, 1H, J=11.2, 6.6Hz), 4.10 (t, 2H, J=7.6Hz), 4.38 (dd, 1H, J=11.2, 10.3Hz), 4.48 (quin, 1H, J=6.6Hz)

IR (CHCl₃): 1734, 1716, 1697, 1683, 1654, 1540 cm⁻¹

EI-MS: m/e 383 (M⁺)

Elemental analysis: for C ₂₂ H ₃₃ N ₅ OúHCl			
Calculated (%):	C, 62.92;	H, 8.16;	N, 16.67

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Elemental analysis: for $C_{22}H_{33}N_5O \cdot HCl$			
Found (%):	C, 62.68;	H, 8.21;	N, 16.50

Example 13: Compound 15

[0059] Compound 15 was obtained from Compound 6a and (R)-phenylglycinol in the same manner as Example 1.

Yield: 49%

Melting point: 215-219°C (chloroform/ether)

Optical rotation: $[\alpha]_D^{20} +87.5^\circ$ {c 1.00, CH_3OH , >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]}

1H NMR (270 MHz, $CDCl_3$) δ : 1.00 (t, 3H, J=7.4Hz), 1.65-2.00 (m, 8H), 2.10-2.25 (m, 2H), 3.20-3.35 (m, 1H), 4.12 (t, 2H, J=7.4Hz), 4.18 (dd, 1H, J=11.5, 7.5Hz), 4.72 (dd, 1H, J=11.5, 10.5Hz), 5.54 (dd, 1H, J=10.0, 7.5Hz), 7.25-7.45 (m, 5H)

IR (KBr): 1726, 1690, 1594 cm^{-1}

EI-MS: m/e 363 (M^+)

Elemental analysis: for $C_{21}H_{25}N_5O \cdot HCl \cdot 0.2H_2O$			
Calculated (%):	C, 62.51;	H, 6.59;	N, 17.33
Found (%):	C, 62.61;	H, 6.99;	N, 16.93

Example 14: Compound 16

[0060] Compound 16 was obtained from Compound 6a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 43%

Melting point: 224-226°C (toluene)

Optical rotation: $[\alpha]_D^{20} +47.3^\circ$ {c 1.00, CH_3OH , >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]}

1H NMR (270 MHz, $CDCl_3$) δ : 0.95 (t, 3H, J=7.4Hz), 1.65-2.00 (m, 8H), 2.15-2.30 (m, 2H), 3.00 (dd, 1H, J=14.0, 8.0Hz), 3.23 (dd, 1H, J=14.0, 4.5Hz), 3.25-3.40 (m, 1H), 4.00-4.15 (m, 3H), 4.21 (dd, 1H, J=9.0, 4.5Hz), 4.65-4.75 (m, 1H), 7.20-7.40 (m, 5H)

IR (KBr): 1722, 1682, 1591 cm^{-1}

EI-MS: m/e 377 (M^+)

Elemental analysis: for $C_{22}H_{27}N_5O \cdot HCl$			
Calculated (%):	C, 63.83;	H, 6.82;	N, 16.92
Found (%):	C, 64.01;	H, 6.88;	N, 16.63

Example 15: Compound 17

[0061] Compound 17 was obtained from Compound 6a and (S)-phenylalaninol in the same manner as Example 1.

Yield: 71%

Melting point: 215-220°C (ethanol)

Optical rotation: $[\alpha]_D^{20}$ -51.3° (c 1.00, CH₃OH, >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

Elemental analysis: for C ₂₂ H ₂₇ N ₅ O ₄ HCl			
Calculated (%):	C, 63.83;	H, 6.82;	N, 16.92
Found (%):	C, 63.68;	H, 7.14;	N, 16.83

Example 16: Compound 18

[0062] Compound 18 was obtained from Compound 6a and 4-fluorophenylalaninol (Indian Journal of Chemistry, Section B, vol. 15B, p.260, 1977) in the same manner as Example 1.

Yield: 47%

Melting point: 226-228°C (methanol/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.4Hz), 1.69-1.91 (m, 8H), 2.12-2.21 (m, 2H), 3.02 (dd, 1H, J=14.5, 7.2Hz), 3.17 (dd, 1H, J=14.5, 4.0Hz), 3.28 (quin, 1H, J=7.9Hz), 4.05 (m, 1H), 4.06 (t, 2H, J=8.6Hz), 4.24 (dd, 1H, J=11.9, 9.6Hz), 4.67 (m, 1H), 7.03 (t, 2H, J=8.6Hz), 7.25 (t, 2H, J=8.6Hz)

IR (CHCl₃) 1717, 1684, 1585, 1512 cm⁻¹

EI-MS: m/e 395 (M⁺)

Elemental analysis: for C ₂₂ H ₂₆ FN ₅ O ₄ HCl			
Calculated (%):	C, 61.18;	H, 6.30;	N, 16.21
Found (%):	C, 61.23;	H, 6.38;	N, 16.16

Example 17: Compound 19

[0063] Compound 19 was obtained from Compound 6a and 3-fluorophenylalaninol (Indian Journal of Chemistry, Section B, vol. 15B, p.260, 1977) in the same manner as Example 1.

Yield: 60%

Melting point: 171-173°C (acetone/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.96 (t, 3H, J=7.4Hz), 1.63-1.94 (m, 8H), 2.08-2.23 (m, 2H), 2.99 (dd, 1H, J=14.2, 7.6Hz), 3.22-3.33 (m, 2H), 4.06 (m, 1H), 4.07 (t, 2H, J=6.3Hz), 4.24 (dd, 1H, J=11.9, 9.9Hz), 4.69 (m, 1H), 6.95-7.08 (m, 3H), 7.31 (m, 1H)

IR (CHCl₃): 1716, 1682, 1585 cm⁻¹

EI-MS: m/e 395 (M⁺)

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Elemental analysis: for $C_{22}H_{26}FN_5O_4HCl$			
Calculated (%):	C, 61.18;	H, 6.30;	N, 16.21
Found (%):	C, 61.03;	H, 6.34;	N, 16.01

5

Example 18: Compound 20

10 [0064] Compound 20 was obtained from Compound 6a and 4-chlorophenylalaninol (DE-A-2206961) in the same manner as Example 1.

Yield: 52%

15

Melting point: 222-224°C (acetone)

1H NMR (270 MHz, $CDCl_3$) δ : 0.95 (t, 3H, J=7.3Hz), 1.64-1.92 (m, 8H), 2.05-2.21 (m, 2H), 3.01 (dd, 1H, J=13.9, 6.9Hz), 3.16 (dd, 1H, J=13.9, 4.6Hz), 3.28 (quin, 1H, J=7.9Hz), 4.03 (dd, 1H, J=11.9, 6.6Hz), 4.06 (t, 2H, J=7.3Hz), 4.24 (dd, 1H, J=11.9, 9.9Hz), 4.66 (m, 1H), 7.22 (d, 2H, J=8.3Hz), 7.31 (d, 2H, J=8.3Hz)

20

IR ($CHCl_3$): 1714, 1682, 1585 cm^{-1}

EI-MS: m/e 411 (M^+)

25

Elemental analysis: for $C_{22}H_{26}ClN_5O_4HCl$			
Calculated (%):	C, 58.93;	H, 6.07;	N, 15.62
Found (%):	C, 58.98;	H, 6.12;	N, 15.45

30

Example 19: Compound 21

35 [0065] Compound 21 was obtained from Compound 6a and 3-chlorophenylalaninol (DE-A-2206961) in the same manner as Example 1.

Yield: 32%

Melting point: 146-148°C (acetone)

40

1H NMR (270 MHz, $CDCl_3$) δ : 0.96 (t, 3H, J=7.6Hz), 1.69-1.90 (m, 8H), 2.15 (m, 2H), 2.98 (dd, 1H, J=14.1, 7.6Hz), 3.20 (dd, 1H, J=14.1, 5.0Hz), 3.28 (quin, 1H, J=7.9Hz), 4.03 (dd, 1H, J=9.9, 6.3Hz), 4.07 (t, 2H, J=6.3Hz), 4.24 (dd, 1H, J=11.4, 9.9Hz), 4.67 (m, 1H), 7.17-7.33 (m, 4H)

45

IR ($CHCl_3$): 1717, 1682, 1585 cm^{-1}

EI-MS: m/e 411 (M^+)

50

Elemental analysis: for $C_{22}H_{26}ClN_5O_4 \cdot 0.9HCl \cdot H_2O$			
Calculated (%):	C, 57.10;	H, 6.29;	N, 15.13
Found (%):	C, 57.01;	H, 6.29;	N, 15.11

55

Example 20: Compound 22

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[0066] Compound 22 was obtained from Compounds 6a and 2,6-dichlorophenylalaninol (DE-A-2206961) in the same manner as Example 1.

Yield: 52%

Melting point: 203-205°C (acetone/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.99 (t, 3H, J=7.6Hz), 1.65-1.86 (m, 8H), 2.15 (m, 2H), 3.27 (quin, 1H, J=7.9Hz), 3.45 (dd, 1H, J=13.8, 8.0Hz), 3.54 (dd, 1H, J=13.8, 6.8Hz), 4.12 (t, 2H, J=7.6Hz), 4.21 (d, 2H, J=7.9Hz), 4.88 (quin, 1H, J=7.9Hz), 7.23 (d, 1H, J=8.9Hz), 7.40 (d, 2H, J=8.6Hz)

IR (CHCl₃): 1715, 1682, 1585 cm⁻¹

El-MS: m/e 445 (M⁺)

Elemental analysis: for C ₂₂ H ₂₅ Cl ₂ N ₅ O ₄ HCl			
Calculated (%):	C, 54.73;	H, 5.43;	N, 14.50
Found (%):	C, 54.68;	H, 5.46;	N, 14.28

Example 21: Compound 23

[0067] Compound 23 was obtained from Compound 6a and 4-bromophenylalaninol (Journal of Medicinal Chemistry, vol. 26 volumes, p.1556, 1983) in the same manner as Example 1.

Yield: 35%

Melting point: 184-186°C (acetone)

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.4Hz), 1.64-1.95 (m, 8H), 2.08-2.24 (m, 2H), 2.99 (dd, 1H, J=13.9, 7.6Hz), 3.15 (dd, 1H, J=13.9, 4.6Hz), 3.25 (m, 1H), 3.99-4.11 (m, 3H), 4.24 (m, 1H), 4.70 (m, 1H), 7.14-7.48 (m, 4H)

IR (CHCl₃): 1717, 1682, 1585 cm⁻¹

El-MS: m/e 455 (M⁺)

Elemental analysis: for C ₂₂ H ₂₆ BrN ₅ O ₄ HCl			
Calculated (%):	C, 53.62;	H, 5.52;	N, 14.21
Found (%):	C, 53.43;	H, 5.53;	N, 13.95

Example 22: Compound 24

[0068] Compound 24 was obtained from Compound 6a and 3-bromophenylalaninol (Journal of Medicinal Chemistry, vol. 26, p.1556, 1983) in the same manner as Example 1.

Yield: 12%

Melting point: 104-106°C (acetone/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.96 (t, 3H, J=7.4Hz), 1.64-1.96 (m, 8H), 2.15 (m, 2H), 2.97 (dd, 1H, J=13.9, 7.6Hz), 3.19 (dd, 1H, J=13.9, 6.2Hz), 3.28 (quin, 1H, J=7.9Hz), 4.03 (dd, 1H, J=11.8, 9.5Hz), 4.07 (t, 2H, J=5.9Hz), 4.23 (dd, 1H, J=12.2, 9.5Hz), 4.68 (m, 1H), 7.21-7.26 (m, 3H), 7.42 (m, 1H)

IR (CHCl₃): 1716, 1682, 1585 cm⁻¹

EI-MS: m/e 455 (M⁺).

5 Example 23: Compound 25

[0069] Compound 25 was obtained from Compound 6a and 4-methoxyphenylalaninol (Tetrahedron Letters, vol. 30, p.4211, 1989) in the same manner as Example 1.

10 Yield: 36%

Melting point: 168-170°C (acetone/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.4Hz), 1.62-1.98 (m, 8H), 2.14-2.23 (m, 2H), 2.98 (dd, 1H, J=11.8, 7.6Hz), 3.14 (dd, 1H, J=11.8, 4.5Hz), 3.28 (quin, 1H, J=7.8Hz), 3.77 (s, 3H), 4.05 (t, 2H, J=8.5Hz), 4.06 (m, 1H), 4.20 (m, 1H), 4.67 (m, 1H), 6.85 (d, 2H, J=7.7Hz), 7.19 (d, 2H, J=7.7Hz)

15 IR (CHCl₃): 1716, 1682, 1585, 1514 cm⁻¹

EI-MS: m/e 407 (M⁺)

Elemental analysis: for C ₂₃ H ₂₉ N ₅ O ₂ ·HCl			
Calculated (%):	C, 62.22;	H, 6.81;	N, 15.77
Found (%):	C, 62.38;	H, 7.02;	N, 15.67

25

Example 24: Compound 26

30 [0070] Compound 26 was obtained from Compound 6a and 4-nitrophenylalaninol (Journal of Chemical Society Perkin Transaction I, p.867, 1989) in the same manner as Example 1.

Yield: 6%

35 Melting point: 215-217°C (acetone)

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.4Hz), 1.62-1.93 (m, 8H), 2.04-2.11 (m, 2H), 3.21-3.38 (m, 3H), 3.98-4.14 (m, 3H), 4.35 (m, 1H), 4.78 (m, 1H), 7.46-7.54 (m, 2H), 8.20-8.25 (m, 2H)

40 IR (CHCl₃): 1716, 1689, 1587, 1525, 1506 cm⁻¹

EI-MS: m/e 422 (M⁺)

Elemental analysis: for C ₂₂ H ₂₆ N ₆ O ₃ ·HCl			
Calculated (%):	C, 57.58;	H, 5.93;	N, 18.31
Found (%):	C, 57.34;	H, 5.91;	N, 17.95

45

50

Example 25: Compound 27

[0071] Compound 27 was obtained from Compound 6a and (R)-O-benzyltyrosinol (Journal of Organic Chemistry, vol. 56, p.1961, 1991) in the same manner as Example 1.

55

Yield: 28%

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Melting point: 202-204°C (acetone/ether)

Optical rotation: $[\alpha]_D^{24}$ -34.5° (c 1.00, CDCl₃, >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.4Hz), 1.62-1.98 (m, 8H), 2.15 (m, 2H), 2.95 (dd, 1H, J=13.8, 7.6Hz), 3.15 (dd, 1H, J=13.8, 4.3Hz), 3.28 (quin, 1H, J=7.6Hz), 4.06 (m, 3H), 4.20 (dd, 1H, J=11.9, 9.5Hz), 4.65 (m, 1H), 5.02 (s, 2H), 6.93 (d, 2H, J=8.6Hz), 7.18 (d, 2H, J=8.6Hz), 7.34-7.41 (m, 5H)

IR (CHCl₃): 1714, 1687, 1682, 1587, 1511 cm⁻¹

EI-MS: m/e 482 (M⁺-H)

Elemental analysis: for C ₂₉ H ₃₃ N ₅ O ₂ ·1.1HCl			
Calculated (%):	C, 66.51;	H, 6.56;	N, 13.37
Found (%):	C, 66.60;	H, 6.60;	N, 13.26

Example 26: Compound 28

[0072] Compound 28 was obtained from Compound 6a and (S)-O-benzyltyrosinol (Journal of Organic Chemistry, vol. 56, p.1961, 1991) in the same manner as Example 1.

Yield: 47%

Melting point: 195-197°C (acetone/ether)

Optical rotation: $[\alpha]_D^{24}$ 31.4° (c 0.72, CDCl₃, >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.5Hz), 1.72-1.88 (m, 8H), 2.15 (m, 2H), 2.95 (dd, 1H, J=13.9, 7.6Hz), 3.15 (dd, 1H, J=13.9, 4.6Hz), 3.27 (quin, 1H, J=7.6Hz), 4.06 (t, 2H, J=7.6Hz), 4.07 (dd, 1H, J=9.9, 2.0Hz), 4.21 (dd, 1H, J=11.8, 9.9Hz), 4.65 (m, 1H), 5.02 (s, 2H), 6.93 (d, 2H, J=8.6Hz), 7.18 (d, 2H, J=8.9Hz), 7.32-7.43 (m, 5H)

IR (CHCl₃): 1716, 1684, 1587, 1512 cm⁻¹

EI-MS: m/e 499 (M⁺)

Elemental analysis: for C ₂₉ H ₃₃ N ₅ O ₃ ·0.6HCl			
Calculated (%):	C, 66.79;	H, 6.49;	N, 13.43
Found (%):	C, 66.61;	H, 6.63;	N, 13.31

Example 27: Compound 29

[0073] Compound 29 was obtained from 8-cyclopentyl-7-methyl-6-methylthio-3-propyl-7H-purin-2(3H)-one (Compound 29a, EP-A-423805) and (R)-phenylalaninol in the same manner as Example 1.

Yield: 12%

Melting point: 142-144°C (ethyl acetate/hexane)

Optical purity: >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]

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¹H NMR (270 MHz, CDCl₃) δ : 0.93 (t, 3H, J=7.4Hz), 1.71 (quin, 2H, J=7.3Hz), 1.81-2.07 (m, 8H), 2.68 (dd, 1H, J=13.5, 8.9Hz), 3.10 (quin, 1H, J=7.9Hz), 3.20 (dd, 1H J=13.5, 5.0Hz), 3.63 (dd, 1H, J=10.9, 6.6Hz), 3.78 (dd, 1H, J=10.9, 9.6Hz), 3.88 (s, 3H), 3.91 (t, 2H, J=7.3Hz), 4.51 (m, 1H), 7.18-7.33 (m, 5H)

IR (CHCl₃): 1717, 1684, 1653, 1439 cm⁻¹

EI-MS: m/e 391 (M⁺).

Example 28: Compound 30

[0074] Compound 30 was obtained from Compound 6a and 2-amino-3,3-diphenyl-1-propanol (Journal of Organic Chemistry, vol. 52, p.1487, 1987) in the same manner as Example 1.

Yield: 15%

Melting point: 178-180°C (ethyl acetate/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.91 (t, 3H, J=7.4Hz), 1.61-1.97 (m, 8H), 2.18 (m, 2H), 3.27 (quin, 1H, J=7.9Hz), 3.76 (d, 1H, J=9.9Hz), 3.97 (dd, 1H, J=11.9, 9.9Hz), 4.18 (t, 2H, J=7.4Hz), 4.55 (d, 1H, J=11.2Hz), 5.21 (q, 1H, J=10.5Hz), 7.10-7.40 (m, 6H), 7.50-7.56 (m, 4H)

IR (CHCl₃): 1715, 1684, 1678, 1599, 1514 cm⁻¹

EI-MS: m/e 453 (M⁺)

Elemental analysis: for C₂₈H₃₁N₅O₂·2.1HCl

Calculated (%):	C, 63.44;	H, 6.29;	N, 13.21
Found (%):	C, 63.58;	H, 6.38;	N, 13.04

Example 29: Compound 31

[0075] Compound 31 was obtained from Compound 6a and 2-amino-4-phenyl-1-butanol (Tetrahedron Letters, vol. 24, p.2935, 1983) in the same manner as Example 1.

Yield: 17%

Melting point: 171-173°C (acetone/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.4Hz), 1.62-1.92 (m, 8H), 1.97-2.23 (m, 4H), 2.79 (t, 1H, J=11.6Hz), 2.81 (t, 1H, J=11.6Hz), 3.22 (m, 1H), 3.85 (m, 1H), 4.02 (t, 2H, J=7.4Hz), 4.27 (m, 2H), 7.20-7.38 (m, 5H)

IR (CHCl₃): 1716, 1682, 1586 cm⁻¹

EI-MS: m/e 391 (M⁺)

Elemental analysis: for C₂₃H₂₉N₅O₂·1.4HCl

Calculated (%):	C, 62.42;	H, 6.92;	N, 15.82
Found (%):	C, 62.40;	H, 6.99;	N, 15.71

Example 30: Compound 32

[0076] 3-Benzyl-8-cyclopentyl-6-methylthio-7H-purin-2(3H)-one (Compound 32a) was obtained by a known method (Journal of Chemical Society Perkin I, p.739, 1973). ¹H NMR (270 MHz, DMSO-d₆) δ : 1.55-2.07 (m, 8H), 2.56 (s, 3H), 3.19 (quin, 1H, J=7.9Hz), 5.19 (s, 2H), 7.21-7.34 (m, 5H), 13.10 (brs, 1H).

[0077] Compound 32 was obtained from Compound 32a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 51%

Melting point: 237-239°C (ethyl acetate/ether)

Optical rotation: [α]_D²⁴ -18.4° {c 0.93, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]}

¹H NMR (270 MHz, CDCl₃) δ : 1.65-2.23 (m, 8H), 2.96 (dd, 1H, J=13.6, 7.6Hz), 3.21 (dd, 1H, J=13.6, 4.3Hz), 3.31 (quin, 1H, J=7.7Hz), 4.05 (dd, 1H, J=9.9, 6.6Hz), 4.18 (dd, 1H, J=11.9, 9.9Hz), 4.67 (m, 1H), 5.24 (s, 2H), 7.24-7.34 (m, 8H), 7.46-7.50 (m, 2H)

IR (CHCl₃): 1716, 1684, 1583 cm⁻¹

EI-MS: m/e 425 (M⁺)

Elemental analysis: for C ₂₆ H ₂₇ N ₅ O ₂ ·0.9HCl			
Calculated (%):	C, 68.13;	H, 6.14;	N, 15.28
Found (%):	C, 68.16;	H, 6.23;	N, 15.19

Example 31: Compound 33

[0078] Compound 33 was obtained from 6-methylthio-3-propyl-7H-purin-2(3H)-one (EP-A-423805, Compound 33a) and (R)-phenylalaninol in the same manner as Example 1.

Yield: 72%

Melting point: 234-236°C (acetone/ether)

Optical purity: >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]}

¹H NMR (270 MHz, CDCl₃) δ : 0.96 (t, 3H, J=7.6Hz), 1.76 (q, 2H, J=7.5Hz), 3.07 (dd, 1H, J=10.6, 7.4Hz), 3.23 (dd, 1H, J=10.8, 4.2Hz), 4.09 (t, 2H, J=7.4Hz), 4.13 (dd, 1H, J=9.8, 7.3Hz), 4.22 (dd, 1H, J=11.8, 9.9Hz), 4.75 (m, 1H), 7.21-7.36 (m, 5H), 7.94 (s, 1H)

IR (CHCl₃): 1717, 1684, 1587 cm⁻¹

EI-MS: m/e 309 (M⁺)

Elemental analysis: for C ₁₇ H ₁₉ N ₅ O ₂ ·2.1HCl			
Calculated (%):	C, 52.91;	H, 5.51;	N, 18.15
Found (%):	C, 53.08;	H, 5.41;	N, 18.26

Example 32: Compound 34

[0079] Compound 33a (5.00 g, 22.3 mmol) was treated with benzyl bromide (2.9 mL) in the presence of sodium

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hydride (60% in oil, 1.78g, 44.5 mmol, 2.0 eq.) in N,N-dimethylformamide (250 ml) to obtain 5.84 g (83%) of 7-benzyl-6-methylthio-3-propyl-7H-purin-2(3H)-one (Compound 34a).

¹H NMR (270 MHz, DMSO-d₆) δ : 0.89 (t, 3H, J=7.3Hz), 1.70 (q, 2H, J=7.3Hz), 2.50 (s, 3H), 4.01 (t, 2H, J=7.6Hz), 5.60 (s, 2H), 7.16 (d, 2H, J=6.6Hz), 7.24-7.39(m, 3H), 8.39 (s, 1H)

[0080] Compound 34 was obtained from Compound 34a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 47%

Melting point: 198-200°C (ethyl acetate/ether)

Optical rotation: [α]_D²⁰ -15.7° (c 0.64, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

¹H NMR (270 MHz, CDCl₃) δ : 0.94 (t, 3H, J=7.6Hz), 1.72 (quin, 2H, J=7.6Hz), 3.17 (dd, 1H, J=13.9, 7.6Hz), 3.38 (dd, 1H, J=13.9, 3.2Hz), 4.00 (dd, 1H, J=11.6, 6.9Hz), 4.02 (t, 2H, J=6.9Hz), 4.12 (dd, 1H, J=11.6, 10.2Hz), 4.88 (m, 1H), 6.00 (d, 1H, J=15.4Hz), 6.20 (d, 1H, J=15.4Hz), 7.16-7.20 (m, 5H), 7.33-7.48 (m, 5H), 7.85 (s, 1H)

IR (CHCl₃): 1716, 1713, 1664, 1579 cm⁻¹

EI-MS: m/e 399 (M⁺)

Elemental analysis: for C ₂₄ H ₂₅ N ₅ O ₄ HCl			
Calculated (%):	C, 66.12;	H, 6.01;	N, 16.06
Found (%):	C, 66.13;	H, 6.08;	N, 16.00

Example 33: Compound 35

[0081] To a solution of 3,8-dipropylxanthine (18.0 g, 76.27 mmol) obtained by a known method (EP-A-038,784) in pyridine (300 mL) was added diphosphorus pentasulfide (35.0 g, 157.66 mmol, 2.0 eq.) and the mixture was heated at 130°C for 2 hours with stirring. The reaction solution was poured into ice water and the mixture was extracted with chloroform. The organic layer was washed with a saturated aqueous solution of sodium chloride and dried, and then the solvent was evaporated. The residue was purified by silica gel column chromatography (chloroform:methanol = 20:1) to obtain a thione derivative (5.4 g, 28%). The thione derivative (5.4 g, 21.43 mmol) was dissolved in a mixture of a 0.5 mol/L aqueous solution of sodium hydroxide (75 mL) and ethanol (30 mL) and the mixture was stirred at room temperature for 30 minutes. To the mixture was added methyl iodide (2 mL) and the mixture was stirred at room temperature for 1 hour. The reaction solution was neutralized with a 2 mol/L solution of hydrochloric acid, and the precipitated crystals were collected by filtration to obtain 6-methylthio-3,8-dipropyl-7H-purin-2(3H)-one (Compound 35a, 3.2 g, 56%).

¹H NMR (270 MHz, CDCl₃) δ : 0.98 (t, 3H, J=7.3Hz), 1.02 (t, 3H, J=7.3Hz), 1.78-1.93 (m, 4H), 2.50 (s, 3H), 2.89 (t, 2H, J=7.3Hz), 4.22 (t, 2H, J=7.6Hz)

[0082] Compound 35 was obtained from Compound 35a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 29%

Melting point: 195-197°C (acetone/ether)

Optical rotation: [α]_D²⁴ -13.6° (c 0.63, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.6Hz), 1.03 (t, 3H, J=6.6Hz), 1.76 (q, 2H, J=7.9Hz), 1.86 (q, 2H,

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J=6.9Hz), 2.84 (dd, 2H, J=8.6, 7.9Hz), 3.00 (dd, 1H, J=13.9, 7.6Hz), 3.23 (dd, 1H, J=13.9, 3.6Hz), 4.05 (t, 2H, J=7.9Hz), 4.06 (m, 1H), 4.21 (t, 1H, J=10.9Hz), 4.70 (m, 1H), 7.23-7.36 (m, 5H)

IR (CHCl₃): 1716, 1506 cm⁻¹

5 EI-MS: m/e 351 (M⁺)

Elemental analysis: for C ₂₀ H ₂₅ N ₅ Oú1.1HCl			
Calculated (%):	C, 61.35;	H, 6.72;	N, 17.89
Found (%):	C, 61.39;	H, 6.80;	N, 17.79

10
15 **Example 34: Compound 36**

[0083] By using 8-cyclopropyl-3-propylxanthine obtained by a known method (EP-A-256,692), the 6-methylthio derivative (Compound 36a) was obtained in the same manner as Example 33.

20 ¹H NMR (270 MHz, DMSO-d₆) δ : 0.85 (t, 3H, J=7.2Hz), 1.00-1.16 (m, 4H), 1.65 (q, 2H, J=7.2Hz), 2.06 (m, 1H), 2.55 (s, 3H), 3.92 (t, 2H, J=6.9Hz)

[0084] Compound 36 was obtained from Compound 36a and (R)-phenylalaninol in the same manner as Example 1.

25 Yield: 26%

Melting point: 187-189°C (acetone/ether)

30 Optical purity: >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]

¹H NMR (270 MHz, CDCl₃) δ : 0.93 (t, 3H, J=7.3Hz), 1.22 (d, 4H, J=6.6Hz), 1.71 (quin, 2H, J=7.3Hz), 2.12 (quin, 1H, J=6.6Hz), 3.00 (dd, 1H, J=13.8, 7.6Hz), 3.22 (dd, 1H, J=13.8, 4.6Hz), 4.01 (t, 2H, J=7.6Hz), 4.06 (dd, 1H, J=11.8, 6.2Hz), 4.20 (dd, 1H, J=11.8, 9.5Hz), 4.68 (m, 1H), 7.24-7.37 (m, 5H)

35 IR (CHCl₃): 1717, 1682, 1585 cm⁻¹

EI-MS: m/e 349 (M⁺)

40 **Example 35: Compound 37**

[0085] By using 8-cyclobutyl-3-propyl-6-thioxanthine obtained by a known method (EP-A-256,692), the 6-methylthio derivative (Compound 37a) was obtained in the same manner as Example 33.

45 ¹H NMR (270 MHz, DMSO-d₆) δ : 0.88 (t, 3H, J=7.2Hz), 1.68 (q, 2H, J=6.9Hz), 1.81-2.10 (m, 2H), 2.21-2.43 (m, 4H), 2.56 (s, 3H), 3.64 (quin, 1H, J=8.9Hz), 3.97 (t, 2H, J=7.4Hz)

[0086] Compound 37 was obtained from Compound 37a and (R)-phenylalaninol in the same manner as Example 1.

50 Yield: 26%

Melting point: 214-216°C (acetone/ether)

55 Optical rotation: [α]_D²⁴ -25.6° (c 1.07, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05])

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¹H NMR (270 MHz, CDCl₃) δ : 0.96 (t, 3H, J=7.6Hz), 1.75 (quin, 2H, J=7.6Hz), 2.03-2.16 (m, 2H), 2.43 (dt, 4H, J=8.6, 6.6Hz), 3.01 (dd, 1H, J=13.8, 7.6Hz), 3.23 (dd, 1H, J=13.6, 4.6Hz), 3.72 (quin, 1H, J=8.6Hz), 4.06 (dd, 1H, J=11.9, 2.9Hz), 4.09 (t, 2H, J=7.5Hz), 4.22 (dd, 1H, J=11.9, 9.6Hz), 4.70 (m, 1H), 7.24-7.37 (m, 5H)

IR (CHCl₃): 1716, 1684, 1587 cm⁻¹

5 EI-MS: m/e 361 (M⁺)

Elemental analysis: for C ₂₁ H ₂₅ N ₅ Oú0.7HClú0.7H ₂ O			
Calculated (%):	C, 62.81;	H, 6.80;	N, 17.44
Found (%):	C, 62.67;	H, 6.80;	N, 17.40

Example 36: Compound 38

[0087] By using 8-cyclohexyl-3-propylxanthine obtained by a known method (Journal of Medicinal Chemistry, vol. 32, p.1231, 1989), the 6-methylthio derivative (Compound 38a) was obtained in the same manner as Example 33.

20 ¹H NMR (270 MHz, CDCl₃) δ : 0.97 (t, 3H, J=7.2Hz), 1.23-1.48 (m, 4H), 1.58 1.96 (m, 6H), 2.01 (s, 3H), 2.06 (d, 2H, J=8.2Hz), 2.87 (m, 1H), 4.25 (t, 2H, J=7.2Hz), 13.08 (brs, 1H)

25 [0088] Compound 38 was obtained from Compound 38a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 28%

Melting point: 243-245øC (acetone/ether)

30 Optical rotation: [α]_D²⁴ -28.2ø {c 1.09, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]}

35 ¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.3Hz), 1.28-1.90 (m, 10H), 2.13 (brd, 2H, J=13.6Hz), 2.87 (tt, 1H, J=11.4, 3.5Hz), 3.00 (dd, 1H, J=13.8, 7.9Hz), 3.24 (dd, 1H, J=13.8, 4.6Hz), 4.05 (dd, 1H, J=11.9, 3.3Hz), 4.08 (t, 2H, J=6.3Hz), 4.21 (dd, 1H, J=11.9, 9.6Hz), 4.70 (m, 1H), 7.21-7.37 (m, 5H)

IR (CHCl₃) 1714, 1636, 1682, 1585 cm⁻¹

40 EI-MS: m/e 391 (M⁺)

Elemental analysis: for C ₂₃ H ₂₉ N ₅ OúHCl			
Calculated (%):	C, 64.55;	H, 7.07;	N, 16.36
Found (%):	C, 64.42;	H, 7.10;	N, 16.19

Example 37: Compound 39

50 [0089] Compound 39 was obtained from Compound 3a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 35%

Melting point: 232-234øC (acetone/ether)

55 Optical rotation: [α]_D²⁴ -41.4ø {c 0.41, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: hexane/ethanol/diethylamine = 95/5/0.05]}

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¹H NMR (270 MHz, CDCl₃) δ : 0.98 (t, 3H, J=7.4Hz), 1.61-2.45 (m, 14H), 2.69 (dd, 1H, J=11.7, 7.3Hz), 2.97 (quin, 1H, J=7.2Hz), 3.21 (dd, 1H, J=11.7, 4.5Hz), 4.07 (dd, 1H, J=11.0, 3.3Hz), 4.08 (t, 2H, J=6.9Hz), 4.20 (t, 1H, J=9.6Hz), 4.66 (m, 1H), 7.23-7.39 (m, 5H)

IR (CHCl₃): 1714, 1693, 1682, 1585 cm⁻¹

EI-MS: m/e 429 (M⁺)

Elemental analysis: for C ₂₆ H ₃₁ N ₅ Oú0.3HClú1.5H ₂ O			
Calculated (%):	C, 66.80;	H, 7.39;	N, 14.98
Found (%):	C, 66.84;	H, 7.09;	N, 15.02

Example 38: Compound 40

[0090] By using 8-cyclopentylmethyl-3-propylxanthine obtained by a known method (Journal of Medicinal Chemistry, vol. 36, p.2508, 1993), the 6-methylthio derivative (Compound 40a) was obtained in the same manner as Example 33.

[0091] Compound 40 was obtained from Compound 40a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 87%

Melting point: 211-213°C (acetone/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.4Hz), 1.21-1.29 (m, 2H), 1.54-1.89 (m, 9H), 2.87 (d, 2H, J=7.6Hz), 3.02 (dd, 1H, J=14.0, 7.8Hz), 3.04 (dd, 1H, J=14.0, 4.6Hz), 4.05 (t, 2H, J=7.1Hz), 4.09 (dd, 1H, J=11.6, 6.6Hz), 4.21 (t, 1H, J=9.9Hz), 4.69 (m, 1H), 7.24-7.36 (m, 5H)

IR (KBr): 1722, 1686, 1676, 1592 cm⁻¹

EI-MS: m/e 391 (M⁺)

Elemental analysis: for C ₂₃ H ₂₉ N ₅ OúHClú0.2H ₂ O			
Calculated (%):	C, 64.01;	H, 7.10;	N, 16.23
Found (%):	C, 63.93;	H, 7.23;	N, 16.56

Example 39: Compound 41

[0092] By using 8-isopropyl-3-propylxanthine obtained by a known method (EP-A-038,784), the 6-methylthio derivative (Compound 41a) was obtained in the same manner as Example 33.

[0093] Compound 41 was obtained from Compound 41a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 48%

Melting point: 169-170°C (acetone/ether)

Optical rotation: [α]_D²⁵ - 17.0° (c 1.15, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 X 250 mm), Daicel Chemical Industries, elution solvent: ethanol])

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.4Hz), 1.42 (d, 6H, J=6.9Hz), 1.77 (q, 2H, J=7.4Hz), 3.00 (dd, 1H, J=13.9, 7.6Hz), 3.20 (q, 1H, J=6.9Hz), 3.21 (dd, 1H, J=13.8, 4.6Hz), 4.07 (t, 2H, J=7.6Hz), 4.08 (dd, 1H, J=12.2,

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5.6Hz), 4.21 (dd, 1H, J=11.9, 9.9Hz), 4.70 (m, 1H), 7.25-7.34 (m, 5H), 11.44 (s, 1H), 13.57 (s, 1H)

IR (KBr): 1729, 1711, 1693, 1678, 1656, 1592, 1540 cm⁻¹

El-MS: m/e 351 (M⁺)

Elemental analysis: for C ₂₀ H ₂₅ N ₅ O ₄ HCl			
Calculated (%):	C, 61.93;	H, 6.76;	N, 18.05
Found (%):	C, 62.14;	H, 6.79;	N, 18.09

Example 40: Compound 42

[0094] 5,6-Diamino-1,2-dihydro-4-mercapto-2-oxo-1-propylpyrimidine (Compound 42a, 1.00 g, 4.99 mmol) obtained by a known method (Journal of Organic Chemistry, vol. 25, p. 1752, 1960) was suspended in a mixed solvent of dioxane (12 mL) and water (6 mL). To the suspension was added 3-furancarboxylic acid (0.56 g, 4.99 mmol) and 1-ethyl-3-[3-(dimethylamino)propyl]carbodiimide hydrochloride (1.63 g, 8.50 mmol) and the mixture was stirred at room temperature for 24 hours. To the reaction mixture was added a 2 mol/L aqueous solution of sodium hydroxide (50 mL), and the mixture was stirred under reflux by heating for 2.5 hours. The mixture was adjusted to pH 6.5 with 4 mol/L hydrochloric acid, and the precipitated solid was collected by filtration. The solid obtained was dissolved in a 0.5 mol/L aqueous solution of sodium hydroxide (7.0 mL). To the solution was added methyl iodide (0.4 mL) and the mixture was stirred at room temperature for 4 hours. The reaction mixture was adjusted to pH 7 with 4 mol/L hydrochloric acid, and the resulting solid was separated by filtration and dried under reduced pressure to obtain the 6-methylthio derivative (Compound 42b, 870 mg, 60%). A mixture of Compound 42b (0.30 g, 1.03 mmol) and (R)-phenylalaninol (0.24 g, 1.55 mmol) in pyridine (4 mL) were heated for 35 hours with stirring, and the reaction solvent was removed under reduced pressure. To the resulting residue was added water and chloroform for extraction, and the organic layer was dried over anhydrous magnesium sulfate. The solvent was evaporated under reduced pressures, and the residue obtained was purified by silica gel column chromatography (chloroform/methanol). The resulting Adduct 42c (0.36 g, 0.92 mmol) was dissolved in chloroform (4 mL). To the solution were added methanesulfonyl chloride (0.142 mL, 1.84 mmol) and pyridine (0.074 mL, 0.92 mmol), the mixture was stirred at room temperature for 20 hours and at 50°C for 7 hours with heating, and then the solvent was evaporated under reduced pressure. To the residue was added water, and the mixture was extracted with chloroform. The organic layer was dried over anhydrous magnesium sulfate and then concentrated. The residue obtained was purified by silica gel column chromatography (chloroform/methanol) and crystallized (hexane/ether) to obtain Compound 42 (0.11 g, 26%).

Melting point: 123-124 °C

Optical rotation: [α]_D²⁴-206.4° {c 1.02, CHCl₃, >99.5% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: ethanol]}

¹H NMR (270 MHz, CDCl₃) δ : 0.97 (t, 3H, J=7.4Hz), 1.75-1.83 (m, 2H), 3.02 (dd, 1H, J=13.9, 7.8Hz), 3.23 (dd, 1H, J=13.9, 4.6Hz), 4.06-4.12 (m, 3H), 4.22 (dd, 1H, J=11.9, 9.9Hz), 4.72-4.73 (m, 1H), 7.10 (dd, 1H, J=1.8, 0.7Hz), 7.28-7.36 (m, 5H), 7.52 (d, 1H, J=1.8Hz), 8.31 (d, 1H, J=0.7Hz)

IR (KBr): 1713, 1709, 1687, 1166, 748 cm⁻¹

El-MS: m/e 375 (M⁺)

Elemental analysis: for C ₂₁ H ₂₁ N ₅ O ₂ ·HCl			
Calculated (%):	C, 61.24;	H, 5.38;	N, 17.00
Found (%):	C, 61.54;	H, 5.28;	N, 17.22

Example 41: Compound 43

[0095] By using 8-tert-butyl-3-propyl-6-thioxanthine, which was obtained from Compound 42a and pivaloyl

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chloride in the same manner as Example 40, the 6-methylthio derivative (Compound 43a) was obtained in the same manner as Example 40. Compound 43 was obtained from Compound 43a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 40%

Melting point: 220-222°C (diethyl ether/ethanol)

Optical rotation: $[\alpha]_D^{25}$ -26.8° (c 1.00, CHCl₃)

¹H-NMR (270 MHz, DMSO-d₆) δ : 0.86 (t, 3H, J=7.5Hz), 1.36 (s, 9H), 1.68 (m, 2H), 3.02 (dd, 1H, J=13.9, 5.9Hz), 3.07 (dd, 1H, J=13.9, 6.3Hz), 3.94 (m, 1H), 3.94 (t, 2H, J=6.9Hz), 4.20 (t, 1H, J=11.2Hz), 4.72 (m, 1H), 7.24-7.32 (m, 5H), 10.3 (brs, 1H), 13.9 (brs, 1H)

IR (CHCl₃): 1714, 1699, 1687, 1673, 1587 cm⁻¹

FAB-MS: m/e 366 (M⁺+1)

Elemental analysis; for C ₂₁ H ₂₇ N ₅ O ₄ HCl			
Calculated (%):	C, 62.75;	H, 7.02;	N, 17.42
Found (%):	C, 62.64;	H, 7.27;	N, 17.47

Example 42: Compound 44

[0096] By using 3-propyl-8-(3-thienyl)-6-thioxanthine obtained in the same manner as Example 40, the 6-methylthio derivative (Compound 44a) was obtained in the same manner as Example 40.

[0097] Compound 44 was obtained from Compound 44a and (R)-phenylalaninol in the same manner as Example 1.

Yield: 39%

Melting point: 196-197°C (ethanol)

Optical rotation: $[\alpha]_D^{24}$ -112.04° (c 0.40, CHCl₃, >99.5% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: ethanol])

¹H NMR (270 MHz, CDCl₃) δ : 0.98 (t, 3H, J=7.4Hz), 1.82 (quin, 2H, J=7.6Hz), 3.04 (dd, 1H, J=13.9, 7.4Hz), 3.23 (dd, 1H, J=13.9, 4.6Hz), 4.08-4.15 (m, 3H), 4.26 (dd, 1H, J=11.9, 9.6Hz), 4.69-4.79 (m, 1H), 7.25-7.38 (m, 5H), 7.45 (dd, 1H, J=5.3, 2.8Hz), 7.74 (dd, 1H, J=5.3, 1.0Hz), 8.22 (dd, 1H, J=2.8, 1.0Hz), 11.1 (brs, 1H), 14.4 (brs, 1H).

IR (KBr): 1712, 1687, 1587 cm⁻¹

EI-MS: m/e 427 (M⁺)

Elemental analysis: for C ₂₁ H ₂₁ N ₅ OS ₂ HCl · 0.2H ₂ O			
Calculated (%):	C, 58.45;	H, 5.23;	N, 16.23
Found (%):	C, 58.44;	H, 5.44;	N, 15.87

Example 43: Compound 45

[0098] Compound 45 was obtained from Compound 38a and 4-bromophenylalaninol in the same manner as Example 1.

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Yield: 51%

Melting point: 210-212°C (ethanol)

¹H-NMR (270 MHz, DMSO-d₆) δ: 0.86 (t, 3H, J=7.3Hz), 1.22-1.80 (m, 10H), 1.97-2.24 (m, 2H), 2.89 (m, 1H), 3.02 (m, 2H), 3.89-3.97 (m, 3H), 4.20 (t, 1H, J=10.9Hz), 4.70 (m, 1H), 7.29 (d, 2H, J=7.9Hz), 7.51 (d, 2H, J=7.9Hz)

IR (neat): 1722, 1716, 1677, 1674, 1587 cm⁻¹

EI-MS: m/e 469 (M⁺)

Elemental analysis: for C ₂₃ H ₂₈ BrN ₅ O ₄ HCl			
Calculated (%):	C, 54.50;	H, 5.77;	N, 13.82
Found (%):	C, 54.56;	H, 6.00;	N, 13.60

Example 44: Compound 46

[0099] Compound 46 was obtained from Compound 35a and 4-chlorophenylalaninol in the same manner as Example 1.

Yield: 63%

Melting point: 191-192°C (ethanol/diisopropyl ether)

¹H NMR (270 MHz, CDCl₃) δ: 0.96 (t, 3H, J=7.3Hz), 1.03 (t, 3H, J=7.3Hz), 1.70-1.80 (m, 2H), 1.81-1.94 (m, 2H), 2.85 (t, 2H, J=7.8Hz), 3.03 (dd, 1H, J=14.2, 6.9Hz), 3.16 (dd, 1H, J=14.2, 4.6Hz), 4.01-4.09 (m, 3H), 4.26 (dd, 1H, J=11.9, 9.9Hz), 4.67-4.69 (m, 1H), 7.23 (d, 2H, J=8.6Hz), 7.31 (d, 2H, J=8.6Hz), 11.5 (brs, 1H), 13.6 (brs, 1H)

IR (KBr): 1716, 1679, 1591 cm⁻¹

EI-MS: m/e 385 (M⁺)

Elemental analysis: for C ₂₀ H ₂₄ ClN ₅ O HCl			
Calculated (%):	C, 56.90;	H, 5.69;	N, 16.60
Found (%):	C, 56.66;	H, 6.07;	N, 16.50

Example 45: Compound 47

[0100] Compound 47 was obtained from Compound 38a and 4-chlorophenylalaninol in the same manner as Example 1.

Yield: 64%

Melting point: 235-236°C (methanol/diisopropyl ether)

¹H NMR (270 MHz, CDCl₃) δ: 0.95 (t, 3H, J=7.4Hz), 1.28-1.89 (m, 8H), 2.10-2.17 (m, 2H), 2.81-2.91 (m, 1H), 3.02 (dd, 1H, J=14.0, 7.1Hz), 3.16 (dd, 1H, J=14.0, 4.8Hz), 4.00-4.09 (m, 3H), 4.24 (dd, 1H, J=11.9, 9.9Hz), 4.65-4.67 (m, 1H), 7.22 (d, 2H, J=8.3Hz), 7.22 (d, 2H, J=8.3Hz), 7.31 (d, 2H, J=8.3Hz), 11.5 (brs, 1H), 13.6 (brs, 1H)

IR (KBr): 1716, 1675, 1587 cm⁻¹

EI-MS: m/e 425 (M⁺)

Elemental analysis: for C ₂₃ H ₂₆ ClN ₅ O ú HCl			
Calculated (%):	C, 59.74;	H, 6.32;	N, 15.14
Found (%):	C, 60.03;	H, 6.38;	N, 15.38

Example 46: Compound 48

[0101] Compound 48 was obtained from Compound 37a and 4-chlorophenylalaninol in the same manner as Example 1.

Yield: 64%

Melting point: 214-215°C (dioxane/diisopropyl ether)

¹H NMR (270 MHz, CDCl₃) δ: 0.96 (t, 3H, J=7.4Hz), 1.71-1.84 (m, 2H), 2.01-2.17 (m, 2H), 2.39-2.49 (m, 4H), 3.02 (dd, 1H, J=14.0, 7.1Hz), 3.16 (dd, 1H, J=14.0, 4.8Hz), 3.65-3.78 (m, 1H), 4.00-4.10 (m, 3H), 4.25 (dd, 1H, J=11.9, 9.9Hz), 4.64-4.72 (m, 1H), 7.22 (d, 2H, J=8.6Hz), 7.32 (d, 2H, J=8.6Hz), 11.5 (brs, 1H), 13.6 (brs, 1H)

IR (KBr): 1716, 1679, 1587 cm⁻¹EI-MS: m/e 397 (M⁺)

Elemental analysis: for C ₂₁ H ₂₄ ClN ₅ O ú HCl			
Calculated (%):	C, 58.07;	H, 5.80;	N, 16.12
Found (%):	C, 57.88;	H, 5.89;	N, 15.93

Example 47: Compound 49

[0102] Compound 49 was obtained from Compound 35a and 4-fluorophenylalaninol in the same manner as Example 1.

Yield: 59%

Melting point: 199-200°C (ethanol/diisopropyl ether)

¹H NMR (270 MHz, CDCl₃) δ: 0.95 (t, 3H, J=7.4Hz), 1.03 (t, 3H, J=7.4Hz), 1.72-1.94 (m, 4H), 2.85 (t, 2H, J=7.8Hz), 2.99-3.18 (m, 2H), 4.03-4.09 (m, 3H), 4.24 (t, 1H, J=10.7Hz), 4.60-4.80 (m, 1H), 7.02 (t, 2H, J=8.4Hz), 7.24-7.28 (m, 2H), 11.5 (brs, 1H)

IR (KBr): 1714, 1575, 1510 cm⁻¹EI-MS: m/e 369 (M⁺)

Elemental analysis: for C ₂₀ H ₂₄ FN ₅ O ú HCl			
Calculated (%):	C, 59.18;	H, 6.21;	N, 17.25
Found (%):	C, 59.14;	H, 6.44;	N, 17.15

Example 48: Compound 50

[0103] Compound 50 was obtained from Compound 38a and 4-fluorophenylalaninol in the same manner as Example 1.

Yield: 46%

Melting point: 237-239°C (ethanol/diisopropyl ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.4Hz), 1.26-1.89 (m, 10H), 2.11-2.15 (m, 2H), 2.86 (tt, 1H, J=11.6, 3.5Hz), 3.02 (dd, 1H, J=14.0, 6.9Hz), 3.17 (dd, 1H, J=14.0, 4.6Hz), 4.02-4.09 (m, 3H), 4.24 (dd, 1H, J=11.9, 9.9Hz), 4.63-4.73 (m, 1H), 7.02 (t, 2H, J=8.6Hz), 7.23-7.28 (m, 2H), 11.4 (brs, 1H), 13.6 (brs, 1H)

IR (KBr): 1718, 1714, 1589, 1510 cm⁻¹

El-MS: m/e 407 (M⁺)

Elemental analysis: for C₂₃H₂₈FN₅O₄HCl

Calculated (%):	C, 61.94;	H, 6.55;	N, 15.70
Found (%):	C, 62.05;	H, 6.83;	N, 15.74

Example 49: Compound 51

[0104] According to a known method, (R)-4-fluorophenylalaninol was obtained from (R)-4-fluorophenylalanine hydrochloride.

[0105] Compound 51 was obtained from Compound 37a and (R)-4-fluorophenylalaninol in the same manner as Example 1.

Yield: 37%

Melting point: 240-245°C (ethyl acetate/acetone)

Optical rotation: [α]_D²⁴ -16.2° {c 1.03, CHCl₃, >99.5% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: ethanol]}

¹H NMR (270 MHz, CDCl₃) δ : 0.96 (t, 3H, J=7.6Hz), 1.78 (quin, 2H, J=7.2Hz), 2.04-2.14 (m, 2H), 2.40-2.50 (m, 4H), 3.02 (dd, 1H, J=13.8, 6.9Hz), 3.17 (dd, 1H, J=12.7, 4.9Hz), 3.72 (quin, 1H, J=8.5Hz), 4.05 (t, 2H, J=5.9Hz), 4.08 (dd, 1H, J=7.6, 2.6Hz), 4.25 (dd, 1H, J=9.9, 2.3Hz), 4.68 (m, 1H), 7.04 (t, 2H, J=8.5Hz), 7.24-7.29 (m, 2H), 11.41 (brs, 1H), 13.70 (brs, 1H)

IR (KBr): 1721, 1712, 1610, 1518 cm⁻¹

El-MS: m/e 381 (M⁺)

Elemental analysis: for C₂₁H₂₄FN₅O₄·HCl

Calculated (%):	C, 60.35;	H, 6.03;	N, 16.76
Found (%):	C, 60.57;	H, 6.22;	N, 16.71

Example 50: Compound 52

[0106] By using 8-butyl-3-propyl-6-thioxanthine obtained in the same manner as Example 40, the 6-methylthio derivative (Compound 52a) was obtained in the same manner as Example 40.

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[0107] Compound 52 was obtained from Compound 52a and 4-fluorophenylalaninol in the same manner as Example 1.

Yield: 20%

Melting point: 193-196°C (acetone/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.2Hz), 0.97 (t, 3H, J=7.2Hz), 1.43 (q, 2H, J=7.6Hz), 1.78 (quin, 4H, J=7.6Hz), 2.87 (t, 2H, J=7.6Hz), 3.02 (dd, 1H, J=14.1, 6.9Hz), 3.16 (dd, 1H, J=14.1, 4.9Hz), 4.05 (dd, 1H, J=12.2, 6.6Hz), 4.06 (t, 2H, J=7.2Hz), 4.24 (dd, 1H, J=11.9, 9.6Hz), 4.66 (m, 1H), 7.03 (t, 2H, J=8.6Hz), 7.23-7.28 (m, 2H), 11.47 (brs, 1H), 13.63 (brs, 1H)

IR (KBr): 1722, 1699, 1678, 1653, 1591 cm⁻¹

EI-MS: m/e 383 (M⁺)

Elemental analysis: for C ₂₁ H ₂₆ N ₅ O ₄ HCl			
Calculated (%):	C, 60.07;	H, 6.48;	N, 16.67
Found (%):	C, 59.98;	H, 6.62;	N, 16.75

Example 51: Compound 53

[0108] Compound 53 was obtained from 8-(2-furyl)-6-methylthio-3-propyl-7H-purin-2(3H)-one (Compound 53a), which was obtained in the same manner as Example 40, and 4-fluorophenylalaninol in the same manner as Example 1.

Yield: 36%

Melting point: 194-195°C (diethyl ether/hexane)

¹H NMR (270 MHz, CDCl₃) δ : 0.97 (t, 3H, J=7.4Hz), 1.66-1.87 (quin, 2H, J=7.5Hz), 3.05 (dd, 1H, J=13.9, 7.1Hz), 3.18 (dd, 1H, J=13.9, 4.5Hz), 4.04-4.14 (m, 3H), 4.26 (dd, 1H, J=11.9, 9.9Hz), 4.68-4.78 (m, 1H), 6.61 (dd, 1H, J=3.6, 2.0Hz), 7.38 (d, 1H, J=3.6Hz), 7.03 (t, 2H, J=8.6Hz), 7.68 (d, 1H, J=2.0Hz), 7.26 (t, 2H, J=8.6Hz), 11.7 (brs, 1H), 14.3 (brs, 1H)

IR (KBr): 1716, 1706, 1700, 1683, 1585 cm⁻¹

EI-MS: m/e 393 (M⁺)

Elemental analysis: for C ₂₁ H ₂₀ N ₅ O ₂ · HCl · 0.8H ₂ O			
Calculated (%):	C, 56.77;	H, 5.13;	N, 15.76
Found (%):	C, 56.78;	H, 5.08;	N, 15.33

Example 52: Compound 54

[0109] Compound 54 was obtained from Compound 42b and (R)-4-fluorophenylalaninol in the same manner as Example 1.

Yield: 35%

Melting point: 149-150°C (diethyl ether/hexane)

Optical rotation: [α]_D²⁴ -349.1° (c 1.03, CHCl₃, >99.5% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: ethanol])

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¹H NMR (270 MHz, CDCl₃) δ : 1.03 (t, 3H, J=7.4Hz), 1.85-1.93 (m, 2H), 2.65 (d, 2H, J=6.6Hz), 3.87-3.90 (m, 1H), 4.10-4.18 (m, 4H), 6.78 (s, 2H), 6.80 (s, 2H), 6.86 (d, 1H, J=1.3Hz), 7.46-7.47 (m, 1H), 8.03 (s, 1H)

IR (KBr): 1695, 1685, 1510, 752 cm⁻¹

5 EI-MS: m/e 393 (M⁺)

Elemental analysis: for C ₂₁ H ₂₀ FN ₅ O ₂			
Calculated (%):	C, 64.11;	H, 5.12;	N, 17.80
Found (%):	C, 64.04;	H, 5.30;	N, 17.78

Example 53: Compound 55

[0110] Compound 55 was obtained from Compound 43a and (R)-4-fluorophenylalaninol in the same manner as Example 1.

20 Yield: 60%

Melting point: 232-236°C (diethyl ether/ethanol)

Optical rotation: [α]_D²⁵ -14.3° {c 1.00, CHCl₃ >99.5% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: ethanol]}

¹H-NMR (270 MHz, DMSO-d₆) δ : 0.86 (t, 3H, J=7.3Hz), 1.36 (s, 9H), 1.68 (m, 2H), 3.04 (m, 2H), 3.95 (m, 1H), 3.95 (t, 2H, J=5.7Hz), 4.21 (t, 1H, J=10.5Hz), 4.69 (m, 1H), 7.13 (dd, 2H, J=8.9, 8.4Hz), 7.36 (dd, 2H, J=8.4, 5.4Hz), 10.4 (brs, 1H), 13.9 (brs, 1H)

30 IR (neat): 1714, 1679, 1583, 1510 cm⁻¹

EI-MS: m/e 384 (M⁺+1)

Elemental analysis: for C ₂₁ H ₂₆ FN ₅ O ₄ HCl			
Calculated (%):	C, 60.06;	H, 6.48;	N, 16.68
Found (%):	C, 60.16;	H, 6.81;	N, 16.80

Example 54: Compound 56

[0111] Compound 56 was obtained from Compound 35a and 3-fluorophenylalaninol in the same manner as Example 1.

45 Yield: 49%

Melting point: 139-140°C (ethanol/diisopropyl ether)

50 ¹H NMR (270 MHz, CDCl₃) δ : 0.96 (t, 3H, J=7.4Hz), 1.03 (t, 3H, J=7.3Hz), 1.73 1.94 (m, 4H), 2.85 (t, 2H, J=7.6Hz), 3.00-3.05 (m, 1H), 3.22-3.27 (m, 1H), 4.04-4.10 (m, 3H), 4.20-4.35 (m, 1H), 4.60-4.83 (m, 1H), 6.96-7.01 (m, 2H), 7.06-7.09 (m, 1H), 7.31-7.36 (m, 1H), 11.5 (brs, 1H)

55 IR (KBr): 1712, 1672, 1585, 1390 cm⁻¹

EI-MS: m/e 369 (M⁺)

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Elemental analysis: for $C_{20}H_{24}FN_5O \cdot HCl \cdot H_2O$			
Calculated (%):	C, 56.67;	H, 6.42;	N, 16.52
Found (%):	C, 56.82;	H, 6.67;	N, 16.48

5

Example 55: Compound 57

10 [0112] Compound 57 was obtained from Compound 37a and 3-fluorophenylalaninol in the same manner as Example 1.

Yield 52%

15

Melting point: 164- 165°C (ethanol/diisopropyl ether)

1H NMR (270 MHz, $CDCl_3$) δ : 0.97 (t, 3H, J=7.4Hz), 1.72-1.85 (m, 2H), 1.99-2.17 (m, 2H), 2.40-2.49 (m, 4H), 3.01 (dd, 1H, J=13.9, 7.6Hz), 3.24 (dd, 1H, J=13.9, 4.6Hz), 3.65-3.78 (m, 1H), 4.02-4.11 (m, 3H), 4.25 (t, 1H, J=10.7Hz), 4.60-4.75 (m, 1H), 6.96-7.08 (m, 3H), 7.29-7.37 (m, 1H), 11.5 (brs, 1H), 13.7 (brs, 1H)

20

IR (KBr): 1722, 1685, 1591, 1515 cm^{-1}

El-MS: m/e 381 (M^+)

25

Elemental analysis: for $C_{21}H_{24}FN_5O \cdot HCl$			
Calculated (%):	C, 60.35;	H, 6.03;	N, 16.76
Found (%):	C, 60.57;	H, 6.22;	N, 16.71

30

Example 56: Compound 58

35 [0113] Compound 58 was obtained from 8-isobutyl-6-methylthio-3-propyl-7H-purin-2(3H)-one (Compound 58a), which was obtained in the same manner as Example 40, and 3-fluorophenylalaninol in the same manner as Example 1.

Yield: 86%

Melting point: 186-187°C (chloroform/diisopropyl ether)

40

1H NMR (270 MHz, $CDCl_3$) δ : 0.89-0.95 (m, 6H), 1.01 (t, 3H, J=7.3Hz), 1.84 (m, 2H), 2.06 (m, 1H), 2.44 (dd, 1H, J=14.2, 7.6Hz), 2.62 (dd, 1H, J=14.2, 7.3Hz), 2.74-2.87 (m, 2H), 3.73 (m, 1H), 3.94-4.08 (m, 4H), 6.75-6.92 (m, 3H), 7.08 (m, 1H)

IR (KBr): 1701, 1653, 1543, 1491 cm^{-1}

45

FAB-MS: m/e 384 ($M^+ + 1$)

50

Elemental analysis: for $C_{21}H_{26}FN_5O$			
Calculated (%):	C, 65.18;	H, 6.83;	N, 18.26
Found (%):	C, 65.80;	H, 7.11;	N, 18.54

55

Example 57: Compound 59

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[0114] Compound 59 was obtained from 8-(4-bromophenyl)-6-methylthio-3-propyl-7H-purin-2(3H)-one (Compound 59a), which was obtained in the same manner as Example 40, and 3-fluorophenylalaninol in the same manner as Example 1.

Yield: 21%

Melting point: 224-226°C (ethyl acetate)

¹H NMR (270 MHz, CDCl₃) δ: 0.99 (t, 3H, J=7.4Hz), 1.81 (m, 2H), 3.05 (q, 1H, J=7.3Hz), 3.21 (dd, 1H, J=11.7, 4.5Hz), 4.07-4.15 (m, 3H), 4.31 (dd, 1H, J=12.2, 10.0Hz), 4.75 (m, 1H), 6.97-7.02 (m, 2H), 7.08 (d, 1H, J=7.6Hz), 7.34 (m, 1H), 7.65 (d, 2H, J=8.5Hz), 8.01 (d, 2H, J=8.5Hz)

IR (KBr): 1706, 1585, 1412, 1351, 1268, 1143, 1070, 1010, 845, 746, 692 cm⁻¹

EI-MS: m/e 481 (M⁺)

Elemental analysis: for C ₂₃ H ₂₁ BrFN ₅ O ₄ HCl _{0.25} CH ₃ CO ₂ C ₂ H ₅			
Calculated (%):	C, 53.30;	H, 4.47;	N, 12.95
Found (%):	C, 53.69;	H, 4.41;	N, 13.01

Example 58: Compound 60

[0115] Compound 60 was obtained from Compound 58a and 2-fluorophenylalaninol (Indian Journal of Chemistry, Section B, vol. 15B, p.260, 1977) in the same manner as Example 1.

Yield: 87%

Melting point: 159-160°C (chloroform/diisopropyl ether)

¹H NMR (270 MHz, CDCl₃) δ: 0.90- 1.03 (m, 9H), 1.82 (m, 2H), 2.12 (m, 1H), 2.51 (dd, 1H, J=14.2, 7.3Hz), 2.67 (dd, 1H, J=14.2, 7.3Hz), 2.78-2.95 (m, 2H), 3.75 (dd, 1H, J=11.2, 5.9Hz), 3.94-4.16 (m, 4H), 6.82 (m, 1H), 6.92 (m, 1H), 7.06-7.11 (m, 2H)

IR (KBr): 1701, 1649, 1545, 1500, 1265 cm⁻¹

FAB-MS: m/e 384 (M⁺+1)

Elemental analysis: for C ₂₁ H ₂₆ FN ₅ O			
Calculated (%):	C, 65.78;	H, 6.83;	N, 18.26
Found (%):	C, 65.83;	H, 7.00;	N, 18.42

[0116] The aminoalcohols used in Examples 59 to 65 were obtained by a known method [Chemical and Pharmaceutical Bulletin, vol. 13 (8), p.995, 1965] from substituted phenylalanine methyl ester hydrochlorides obtained by a known method.

Example 59: Compound 61

[0117] Compound 61 was obtained from Compound 6a and 3-methylphenylalaninol in the same manner as Example 1.

Yield: 50%

Melting point: 157-159°C (ethyl acetate/diisopropyl ether)

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¹H NMR (270 MHz, CDCl₃) δ : 0.97 (t, 3H, J=7.4Hz), 1.67-2.09 (m, 10H), 2.23 (s, 3H), 2.82 (dd, 1H, J=13.5, 6.9Hz), 2.97 (dd, 1H, J=13.5, 6.6Hz), 3.15-3.21 (m, 1H), 3.85 (dd, 1H, J=11.2, 6.6Hz), 4.02-4.09 (m, 3H), 4.25-4.31 (m, 1H), 6.89 (brs, 1H), 6.96-7.26 (m, 4H)

IR (KBr): 1702, 1681, 1542 cm⁻¹

EI-MS: m/e 391 (M⁺)

Elemental analysis: for C ₂₃ H ₂₉ N ₅ O ú 0.5H ₂ O			
Calculated (%):	C, 68.97;	H, 7.55;	N, 17.49
Found (%):	C, 69.30;	H, 7.70;	N, 17.42

Example 60: Compound 62

[0118] Compound 62 was obtained from Compound 6a and 3-iodophenylalaninol in the same manner as Example 1.

Yield: 35%

Melting point: 146-148°C (acetone/ether)

¹H NMR (270 MHz, CDCl₃) δ : 0.96 (t, 3H, J=7.4Hz), 1.70-1.89 (m, 8H), 2.13-2.23 (m, 2H), 2.95 (dd, 1H, J=13.0, 7.3Hz), 3.15 (dd, 1H, J=13.0, 4.8Hz), 3.28 (quin, 1H, J=7.9Hz), 4.04 (dd, 1H, J=8.8, 2.9Hz), 4.07 (t, 2H, J=6.6Hz), 4.23 (t, 1H, J=9.7Hz), 4.68 (m, 1H), 7.08 (t, 1H, J=7.9Hz), 7.24 (d, 1H, J=1.0Hz), 7.61 (dd, 2H, J=7.9, 1.0Hz)

IR (KBr): 1714, 1684, 1670, 1587 cm⁻¹

EI-MS: m/e 503 (M⁺)

Elemental analysis: for C ₂₂ H ₂₆ N ₅ OI ú HCl			
Calculated (%):	C, 48.95;	H, 5.04;	N, 12.97
Found (%):	C, 48.87;	H, 5.14;	N, 12.75

Example 61: Compound 63

[0119] Compound 63 was obtained from Compound 6a and 2,3-difluorophenylalaninol in the same manner as Example 1.

Yield: 53%

Melting point: 185-187°C (diisopropyl ether/ethanol)

¹H-NMR (270 MHz, DMSO-d₆) δ : 0.90 (t, 3H, J=7.6Hz), 1.65-1.77 (m, 8H), 2.08 (m, 2H), 3.17 (m, 2H), 3.29 (m, 1H), 3.96 (t, 2H, J=11.2Hz), 3.96 (m, 1H), 4.30 (t, 1H, J=10.6Hz), 4.73 (m, 1H), 7.16-7.41 (m, 3H), 10.6 (brs, 1H), 13.8 (brs, 1H)

IR (neat): 1714, 1682, 1587, 1489 cm⁻¹

FAB-MS: m/e 414 (M⁺+1)

Elemental analysis: for C ₂₂ H ₂₅ F ₂ N ₅ O ú HCl ú 0.3H ₂ O			
Calculated (%):	C, 58.03;	H, 5.89;	N, 15.38

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Elemental analysis: for $C_{22}H_{25}F_2N_5O \cdot HCl \cdot 0.3H_2O$			
Found (%)	C, 57.92;	H, 5.88;	N, 15.60

Example 62: Compound 64

[0120] Compound 64 was obtained from Compound 6a and 2,5-difluorophenylalaninol in the same manner as Example 1.

Yield: 35%

Melting point: 198-202°C (acetone/ether)

1H NMR (270 MHz, $CDCl_3$) δ : 0.97 (t, 3H, $J=7.5$ Hz), 1.67-1.92 (m, 8H), 2.04-2.23 (m, 2H), 3.04 (dd, 1H, $J=13.6, 7.2$ Hz), 3.21-3.36 (m, 2H), 3.98-4.14 (m, 3H), 4.27 (dd, 1H, $J=12.2, 9.6$ Hz), 4.73 (m, 1H), 6.93-7.18 (m, 3H), 11.56 (brs, 1H), 13.44 (brs, 1H)

IR (KBr): 1716, 1684, 1652, 1585, 1506, 1498 cm^{-1}

EI-MS: m/e 413 (M^+)

Elemental analysis: for $C_{22}H_{25}F_2N_5O \cdot HCl \cdot 0.1H_2O$			
Calculated (%)	C, 58.50;	H, 5.85;	N, 15.50
Found (%)	C, 58.57;	H, 5.86;	N, 15.43

Example 63: Compound 65

[0121] Compound 65 was obtained from Compound 6a and 2,5-dichlorophenylalaninol in the same manner as Example 1.

Yield: 42%

Melting point: 172-173°C (ethyl acetate/ether)

1H NMR (270 MHz, $CDCl_3$) δ : 0.98 (t, 3H, $J=7.4$ Hz), 1.67-1.94 (m, 8H), 2.12-2.22 (m, 2H), 3.12 (dd, 1H, $J=14.2, 7.8$ Hz), 3.25-3.31 (m, 1H), 3.37 (dd, 1H, $J=14.2, 5.6$ Hz), 4.05-4.13 (m, 3H), 4.26 (dd, 1H, $J=11.9, 9.9$ Hz), 4.76-4.82 (m, 1H), 7.22-7.28 (m, 1H), 7.34-7.38 (m, 2H), 11.5 (brs, 1H)

IR (KBr): 1714, 1668, 1585 cm^{-1}

EI-MS: m/e 445 (M^+)

Elemental analysis: for $C_{22}H_{25}Cl_2N_5O \cdot HCl$			
Calculated (%)	C, 54.72;	H, 5.18;	N, 14.50
Found (%)	C, 55.03;	H, 5.27;	N, 14.61

Example 64: Compound 66

[0122] Compound 66 was obtained from Compound 3a and (S)-tert-leucinol in the same manner as Example 1.

Yield: 27%

Melting point: 230-233°C (acetone/ether)

Optical purity ; >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: ethanol]

¹H NMR (270 MHz, CDCl₃) δ : 0.98 (t, 3H, J=7.6Hz), 1.06 (s, 9H), 1.62-1.73 (m, 3H), 1.82 (q, 2H, J=7.3Hz), 1.94-2.02 (m, 3H), 2.03 (d, 2H, J=10.9Hz), 2.24 (d, 2H, J=10.4Hz), 2.41 (brs, 2H), 2.70 (t, 1H, J=6.6Hz), 4.12 (t, 2H, J=7.3Hz), 4.14-4.26 (m, 3H), 11.29 (brs, 1H), 13.54 (brs, 1H)
IR (CHCl₃) 1816, 1794, 1714, 1683, 1652, 1585 cm⁻¹EI-MS: m/e 395 (M⁺)

Elemental analysis: for C ₂₃ H ₃₃ N ₅ O ₄ HCl·0.2H ₂ O			
Calculated (%):	C, 63.42;	H, 7.96;	N, 16.08
Found (%):	C, 63.52;	H, 8.16;	N, 15.94

Example 65: Compound 67

[0123] Compound 67 was obtained from Compound 42b and (S)-tert-leucinol in the same manner as Example 40.

Yield: 38%

Melting point: 204-206°C (ethyl acetate/hexane)

Optical rotation: [α]_D²⁴ 253.9° (c 1.03, CHCl₃, >99% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel Chemical Industries, elution solvent: ethanol])
¹H NMR (270 MHz, CDCl₃) δ : 0.77 (s, 9H), 1.01 (t, 3H, J=7.6Hz), 1.86 (quin, 2H, J=7.3Hz), 3.87 (dd, 1H, J=10.5, 6.3Hz), 4.03 (dd, 1H, J=12.2, 6.1Hz), 4.16 (m, 3H), 6.89 (s, 1H), 7.45 (s, 1H), 8.06 (s, 1H)
IR (KBr): 1689, 1683, 1548, 1514, 1371, 1306 cm⁻¹EI-MS: m/e 341 (M⁺)

Elemental analysis: for C ₁₈ H ₂₃ N ₅ O ₂ · H ₂ O			
Calculated (%):	C, 60.15;	H, 7.01;	N, 19.48
Found (%):	C, 60.24;	H, 7.28;	N, 19.49

Example 66: Compound 68

[0124] Compound 68 was obtained from 8-(3,5-dimethylisoxazol-4-yl)-6-methylthio-3-propyl-7H-purin-2(3H)-one (Compound 68a), which was obtained in the same manner as Example 40, and (S)-tert-leucinol in the same manner as Example 1.

Yield: 63%

Melting point: 185-186°C (ethyl acetate/hexane)

Optical rotation: [α]_D²⁵ -13.68° (c 1.01, CHCl₃, >99.5% ee [CHIRALCEL OD column (4.6 x 250 mm), Daicel

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Chemical Industries, elution solvent: ethanol]]

¹H NMR (270 MHz, CDCl₃) δ: 1.01 (t, 3H, J=7.4Hz), 1.09 (s, 9H), 1.79-1.93 (m, 2H), 2.67 (s, 3H), 2.87 (s, 3H), 4.09-4.32 (m, 5H), 11.8 (brs, 1H), 13.9 (brs, 1H)

IR (KBr): 1714, 1685, 1413 cm⁻¹

FAB-MS: m/e 371 (M⁺+1)

Example 67: Compound 69

[0125] Compound 69 was obtained from Compound 41a and (S)-valinol in the same manner as Example 1.

Yield: 55%

Melting point: 156- 157°C (dioxane/ether)

Optical rotation: [α]_D²⁵ -44.14° (c 1.01, CHCl₃, >99.5% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: ethanol]]

¹H NMR (270 MHz, CDCl₃) δ: 0.99 (t, 3H, J=7.4Hz), 1.06 (d, 3H, J=6.9Hz), 1.15 (d, 3H, J=6.6Hz), 1.42 (d, 6H, J=6.9Hz), 1.64-1.86 (m, 2H), 1.93-2.01 (m, 1H), 3.17 (quin, 1H, J=6.9Hz), 4.01 (dd, 1H, J=11.1, 7.1Hz), 4.08-4.22 (m, 3H), 4.33 (dd, 1H, J=11.1, 10.1Hz), 11.5 (brs, 1H), 13.6 (brs, 1H)

IR (KBr): 1712, 1687, 1592 cm⁻¹

El-MS: m/e 304 (M⁺)

Elemental analysis: for C ₁₆ H ₂₅ N ₅ O · HCl			
Calculated (%):	C, 56.54;	H, 7.71;	N, 20.61
Found (%):	C, 56.69;	H, 7.95;	N, 20.61

Example 68: Compound 70

[0126] 3-Cyanophenylalanine methyl ester hydrochloride disclosed in DE 19544687 was converted into an amino alcohol by a known method [Chemical and Pharmaceutical Bulletin, vol. 13 (8), p.995, 1965]. By using this compound and Compound 6a, Compound 70 was obtained in the same manner as Example 1.

Yield: 58%

¹H-NMR (270 MHz, CDCl₃) δ: 0.99 (t, 3H, J=7.3Hz), 1.63-1.94 (m, 10H), 2.80 (m, 1H), 2.95 (dd, 1H, J=13.5, 5.3Hz), 3.04 (dd, 1H, J=13.5, 8.9Hz), 3.80 (dd, 1H, J=11.2, 6.3Hz), 4.00 (t, 2H, J=7.3Hz), 4.10 (t, 1H, J=11.2Hz), 4.37 (m, 1H), 7.30-7.34 (m, 2H), 7.47 (d, 1H, J=6.9Hz), 7.57 (s, 1H)

Example 69: Compound 71

[0127] Compound 71 was obtained from 8-cyclopentyl-3-ethyl-6-methylthio- 7H-purin-2(3H)-one (Compound 71a), which was obtained in the same manner as Example 40, and (R)-phenylalaninol in the same manner as Example 1.

Yield: 44%

Melting point: 168-169°C (dioxane/ether)

Optical rotation: [α]_D²⁵ -30.40° (c 1.02, CHCl₃, >99.5% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: ethanol]]

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¹H NMR (270 MHz, CDCl₃) δ: 1.33 (t, 3H, J=7.1Hz), 1.70-2.23 (m, 8H), 2.99 (dd, 1H, J=14.0, 7.6Hz), 3.23 (dd, 1H, J=14.0, 4.6Hz), 3.21-3.31 (m, 1H), 4.08 (dd, 1H, J=12.2, 6.6Hz), 4.13-4.25 (m, 3H), 4.65-4.80 (m, 1H), 7.25-7.37 (m, 5H), 11.4 (brs, 1H), 13.7 (brs, 1H)

IR (KBr): 1720, 1712, 1679, 1591 cm⁻¹

FAB-MS: m/e 364 (M⁺+1)

Elemental analysis: for C ₂₁ H ₂₅ N ₅ O · HCl · 0.2H ₂ O			
Calculated (%):	C, 62.50;	H, 6.59;	N, 17.36
Found (%):	C, 62.65;	H, 6.67;	N, 17.36

Example 70: Compound 72

[0128] In the same manner as Example 1, Compound 72 was obtained from (R)-phenylalaninol and 8-cyclopentyl-3-cyclopropylmethyl-6-methylthio-7H-purin-2(3H)-one (Compound 72a), which was obtained by a known method (Journal of Medicinal Chemistry, vol. 36, p.2508, 1993) using a known compound, 5,6-diamino-1-cyclopropylmethyl-2,4-pyrimidinedione, disclosed in EP-A- 386683, and (R)-phenylalaninol.

Yield: 56%

¹H NMR (270 MHz, CDCl₃) δ: 0.44-0.56 (m, 4H), 1.23-1.37 (m, 1H), 1.70-2.19 (m, 8H), 2.96 (dd, 1H, J=13.7, 7.8Hz), 3.16-3.31 (m, 2H), 3.97 (d, 2H, J=7.3Hz), 3.96-4.04 (m, 1H), 4.16 (t, 1H, J=10.7Hz), 4.50-4.68 (m, 1H), 7.23-7.45 (m, 5H), 9.60 (brs, 1H) FAB-MS: m/e 390 (M⁺+1)

Example 71: Compound 73

[0129] Compound 73 was obtained from 8-cyclopropylmethyl-6-methylthio-3-propyl-7H-purino-2(3H)-one (Compound 73a), which was obtained in the same manner as Example 40, and (R)-phenylalaninol in the same manner as Example 1.

Yield: 56%

Melting point: 188-190°C (ether/ethanol)

Optical rotation: [α]_D²⁵ -22.0° (c 1.03, CHCl₃)

¹H-NMR (270 MHz, DMSO-d₆) δ: 0.28 (m, 2H), 0.56 (m, 2H), 0.87 (t, 3H, J=7.3Hz), 1.05 (m, 1H), 1.66 (m, 2H), 2.78 (d, 2H, J=6.6Hz), 3.01 (dd, 1H, J=13.2, 6.3Hz), 3.19 (dd, 1H, J=13.2, 6.3Hz), 3.95 (m, 1H), 3.95 (t, 2H, J=6.9Hz), 4.20 (t, 1H, J=10.9Hz), 4.74 (m, 1H), 7.21-7.33 (m, 5H), 10.7 (brs, 1H), 13.7 (brs, 1H)

IR (neat): 1713, 1682, 1673, 1587 cm⁻¹

FAB-MS: m/e 364 (M⁺+1)

Elemental analysis: for C ₂₁ H ₂₅ N ₅ O · HCl			
Calculated (%):	C, 63.07;	H, 6.55;	N, 17.51
Found (%):	C, 63.04;	H, 6.85;	N, 17.47

Example 72: Compound 74

[0130] To Compound 70 (100 mg, 0.25 mmol) were added a 4 mol/L aqueous solution of sodium hydroxide (2 mL)

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and methanol (2 mL), and the mixture was stirred at 100°C for 6 hours. The reaction mixture was neutralized with 2 mol/L hydrochloric acid, and then the solid precipitated was collected by filtration to obtain Compound 74 (84 mg, 80%) as a white solid.

Yield: 80%

Melting point: >300°C

¹H-NMR (270 MHz, DMSO-d₆) δ : 0.84 (t, 3H, J=7.3Hz), 1.58-1.72 (m, 8H), 1.94 (m, 2H), 2.95 (m, 2H), 3.07 (q, 1H, J=9.5Hz), 3.59 (dd, 1H, J=11.3, 7.0Hz), 3.83 (t, 2H, J=6.5Hz), 3.92 (t, 1H, J=10.5Hz), 4.50 (m, 1H), 7.34 (t, 1H, J=8.0Hz), 7.50 (d, 1H, J=8.0Hz), 7.76 (d, 1H, J=8.0Hz), 7.84 (s, 1H)

IR (neat): 1713, 1674, 1662 cm⁻¹

EI-MS: m/e 421 (M⁺)

Example 73: Compound 75

[0131] β,β -Dimethylphenylalanine obtained by a known method [Acta Pharmaceutica Suecica, vol. 13, p.75, 1976] was converted into 2-amino-3-methyl-3-phenyl-1-butanol by a known method. From this compound and Compound 6a, Compound 75 was obtained in the same manner as Example 1.

Yield: 25%

Melting point: 85-90°C

¹H-NMR (270 MHz, CDCl₃) δ : 0.94 (t, 3H, J=7.3Hz), 1.42 (s, 3H), 1.45 (s, 3H), 1.68-1.85 (m, 8H), 2.10 (m, 2H), 3.21 (q, 1H, J=7.8Hz), 3.81 (dd, 1H, J=6.8, 11.9Hz), 3.88 (t, 1H, J=11.9Hz), 4.00 (t, 2H, J=7.3Hz), 4.41 (dd, 1H, J=6.8, 9.5Hz), 7.16-7.38 (m, 5H)

IR (neat): 1716, 1704, 1693, 1660 cm⁻¹

EI-MS: m/e 405 (M⁺)

Example 74: Compound 76

[0132] To a solution of Compound 27 (91.2 mg, 0.176 mmol) in methanol (2.0 mL) were added 20% Pd(OH)₂/C (15 mg) and ammonium formate (90 mg) and the mixture was refluxed for 2 hours. The reaction mixture was filtered and concentrated, to the residue was added water and the mixture was extracted with ethyl acetate. Crystallization was performed from acetone/ether to obtain Compound 76 (41.2 mg, 60%).

Yield: 60%

Melting point: 148-151°C (acetone/ether)

Optical purity: >99% ee [CHIRALCEL OD column (4.6 × 250 mm), Daicel Chemical Industries, elution solvent: ethanol]

¹H NMR (270 MHz, CDCl₃) δ : 0.95 (t, 3H, J=7.3Hz), 1.63-1.98 (m, 8H), 2.04-2.18 (m, 2H), 2.90 (dd, 1H, J=14.2, 7.6Hz), 3.07 (dd, 1H, J=13.9, 4.3Hz), 3.24 (quin, 1H, J=8.2Hz), 4.03 (dd, 1H, J=11.8, 6.3Hz), 4.05 (t, 2H, J=7.6Hz), 4.15 (dd, 1H, J=11.9, 9.9Hz), 4.60 (m, 1H), 6.77 (d, 2H, J=8.3Hz), 7.03 (d, 2H, J=8.3Hz), 8.62 (s, 1H)

IR (CHCl₃) 1716, 1682, 1581, 1558, 1540, 1385 cm⁻¹

EI-MS: m/e 393 (M⁺)

Example 75: Compound 77

[0133] To a solution of 6-methylthio-3-propyl-8-(3-pyridyl)-7H-purin-2(3H)-one (15 mg, 50 μmol) obtained by a

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known method (Journal of Medicinal Chemistry, vol. 36, p. 2508, 1993) or in the same manner as Example 40 in chloroform or chloroform/methanol (4/1, 0.20 mL) was added a solution of (R)-phenylalaninol (0.50 mmol/L) in methanol (0.20 mL) and the mixture was heated at 80°C with stirring. After the solvent was removed, the reaction mixture was stirred at 150°C for 5 hours. To the reaction residue were added dichloromethane (2.0 mL), pyridine (two drops) and methanesulfonyl chloride (two drops), and the mixture was stirred at room temperature for 3 hours. The reaction solvent was evaporated, and to the residue were added water (1.0 mL) and ethyl acetate (1.0 mL). The mixture was passed through a diatomaceous earth column, the column was washed with ethyl acetate (6.0 mL), and then the eluates were concentrated. A solution of the residue in dichloromethane (1.0 mL) was applied to a solid phase extraction cartridge (SCX, 500 mg), and the cartridge was washed with dichloromethane (20 mL) and methanol (3.0 mL). Compound 77 was eluted with a 2 mol/L ammonia solution in methanol (3 mL) and the eluate was concentrated. The residue was purified by silica gel column chromatography to obtain Compound 77 (7.6 mg, 39%).

ESI-MS: m/e 387 (M⁺+1)

[0134] Compounds 78 to 92 were obtained in the same manner as Example 75.

[0135] The aminoalcohols used in Examples 84 to 90 (Table 2) were prepared as follows. To a solution of diphenyliminoglycine (53.4 mg, 200 mmol) in N,N-dimethylformamide (0.8 mL) were added 2-tert-butylimino-2-diethylamino-1,3-dimethylperhydro-1,3,2-diazaphospholine polystyrene resin (< 2.3 mmol base/g, 0.2 g) and a corresponding substituted benzyl halide (300 mmol), and the mixture was stirred overnight at room temperature. The reaction mixture was filtered and concentrated under reduced pressure. To the residue were added tetrahydrofuran (800 mL) and 1 mol/L hydrochloric acid (100 mL), the mixture was stirred at room temperature for 2 hours, and then the solvent was evaporated under reduced pressure. To the residue were added 50% ethanol/water (600 mL) and a solution of sodium borohydride (76 mg, 2 mmol) in 50% ethanol/water (2 mL), and the mixture was stirred at room temperature for two days. The solvent was removed under reduced pressure, to the resulting residue were added ethyl acetate (3 mL) and water (0.1 mL), and the mixture was passed through a diatomaceous earth column. Ethyl acetate was applied (3 mL x 2), the resulting eluate was concentrated under reduced pressure, and the residue was treated in a solid phase extraction cartridge (SCX, 500 mg) to obtain a substituted phenylalaninol in an approximate yield of 82 to 100%.

Table 2

Example	Substituted benzyl halide	Aminoalcohol
84	4-tert-Butylbenzyl bromide	2-Amino-3-(4-tert-butylphenyl)-1-propanol
85	2-Phenylbenzyl bromide	2-Amino-3-(2-biphenyl)-1-propanol
86	3-Trifluoromethylbenzyl bromide	2-Amino-3-(3-trifluoromethylphenyl)-1-propanol
87	3-(2-Fluorophenoxy)benzyl bromide	2-Amino-3-[3-(2-fluorophenoxy)phenyl]-1-propanol
88	3-Phenoxybenzyl chloride	2-Amino-3-(3-phenoxyphenyl)-1-propanol
89	3-Trifluoromethoxybenzyl bromide	2-Amino-3-(3-trifluoromethoxyphenyl)-1-propanol
90	3,5-Bis(trifluoromethyl)benzyl bromide	2-Amino-3-[3,5-bis(trifluoromethyl) phenyl]-1-propanol

Example 76: Compound 78

[0136]

EI-MS: m/e 386 (M⁺)

Example 77: Compound 79

[0137]

ESI-MS: m/e 406 (M⁺+1)

Example 78: Compound 80

[0138]

ESI-MS: m/e 430 (M⁺+1)

Example 79: Compound 81

[0139]

5 ESI-MS: m/e 387 ($M^{+}+1$)

Example 80: Compound 82

10 [0140]

ESI-MS: m/e 388 ($M^{+}+1$)

Example 81: Compound 83

15 [0141]

ESI-MS: m/e 392 ($M^{+}+1$)

20

Example 82: Compound 84

[0142]

25 ESI-MS: m/e 350 ($M^{+}+1$)

Example 83: Compound 85

30 [0143]

ESI-MS: m/e 412 ($M^{+}+1$)

Example 84: Compound 86

35

[0144]

EI-MS: m/e 433 (M^{+})

40

Example 85: Compound 87

[0145]

45 EI-MS: m/e 453 (M^{+})

Example 86: Compound 88

[0146]

50

EI-MS: m/e 445 (M^{+})

Example 87: Compound 89

55

[0147]

EI-MS: m/e 487 (M^+)

Example 88: Compound 90

[0148]

EI-MS: m/e 469 (M^+)

Example 89: Compound 91

[0149]

EI-MS: m/e 461 (M^+)

Example 90: Compound 92

[0150]

EI-MS: m/e 513 (M^+)

Example 91: Compound 93

[0151] Compound 93 was obtained from 2-amino-2-methyl-3-phenyl-1-propanol, which was obtained from α -methylphenylalanine by a known method, and Compound 6a in the same manner as Example 1.

Yield: 10%

Melting point: 130-135°C

1H -NMR (270 MHz, $CDCl_3$) δ : 0.94 (t, 3H, $J=7.3$ Hz), 1.42 (s, 3H), 1.45 (s, 3H), 1.68-1.85 (m, 8H), 2.10 (m, 2H), 3.21 (q, 1H, $J=7.8$ Hz), 3.81 (dd, 1H, $J=11.9, 6.8$ Hz), 3.88 (t, 1H, $J=11.9$ Hz), 4.00 (t, 2H, $J=7.3$ Hz), 4.41 (dd, 1H, $J=9.5, 6.8$ Hz), 7.16-7.38 (m, 5H)

IR (neat): 1695, 1651, 1495 cm^{-1}

EI-MS: m/e 391 (M^+)

Test Example 1: Acute toxicity test

[0152] A test compound was orally administered to dd male mice (body weight: 20 \pm 1 g, $n=3$). Mortality rate after seven days was measured to determine minimum lethal dose (MLD). As a result, MLD of Compound 16 was not less than 300 mg/kg (mice, po), which revealed safety of the medicament of the present invention.

Test Example 2: Insulin secretion enhancing action in cultured β -cell

[0153] The established pancreas β -cell, MIN6 cell, reported by Miyazaki et al. (Endocrinology, vol. 127, pp.126-131, 1990) exhibits insulin content and insulin secretion amount by stimulation with glucose similar to those of pancreas β -cells in living bodies, and well preserves characteristics of pancreas β -cells in living bodies from a viewpoint that it shows increase of insulin secretion in a glucose concentration-dependent manner (the above reference and Ishihara et al., Diabetologia, vol. 36, pp.1139-1145, 1993). Further, the insulin secretion of the MIN6 cell is enhanced in response to sulfonylurea agents such as glibenclamide, which are used as a medicament for treatment of diabetes (Beng et al., Cellular Signalling, vol. 5, pp.777-786, 1993).

[0154] Culture of the MIN6 cells and insulin secretion test utilizing the MIN6 cells were performed basically according to the methods described in literature (Ishihara et al., Diabetologia, vol. 36, pp.1139-1145, 1993). The effect of a compound on the insulin secretion in the presence of 14.5 mM glucose was determined by measuring insulin amounts in cell culture supernatants collected as follows. MIN6 cells cultured on a 24-well plate were washed twice by using 1 ml of Buffer A (119 mM sodium chloride, 4.74 mM potassium chloride, 2.54 mM calcium chloride, 1.19 mM magnesium sulfate, 1.19 mM potassium dihydrogenphosphate, 10 mM 2-[4-(2-hydroxyethyl)-1-

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piperaziny]ethanesulfonic acid, 0.1% bovine serum albumin, pH 7.3) containing 2 mM glucose, and then were incubated in 1 ml of Buffer A containing 2 mM glucose at 37°C for 45 minutes. After the incubation, the culture supernatant was changed to Buffer A (0.9 ml) containing a test compound at various concentrations and 2 mM glucose, and the cells were further incubated at 37°C for 15 minutes. The MIN6 cells were stimulated with glucose by the addition of Buffer A (0.1 ml) containing 127 mM glucose to the culture (final glucose concentration: 14.5 mM). After the stimulation, the cells were further incubated at 37°C for 45 minutes, and then the culture supernatant was collected.

[0155] Separately, the effect of a compound on the insulin secretion in the presence of 5 mM glucose was determined by measuring insulin amounts in cell culture supernatants collected as follows. MIN6 cells cultured on a 24-well plate were washed twice by using 1 ml of Buffer A containing 5 mM glucose, and then the culture supernatant was changed to Buffer A (0.9 ml) containing a test compound at various concentrations and 5 mM glucose. Then, the cells were incubated at 37°C for 45 minutes (final glucose concentration: 5 mM), and the culture supernatant was collected.

[0156] After the culture supernatant was diluted with a phosphate buffer containing 1% bovine serum albumin, 0.1% Tween 20, 0.12% disodium ethylenediaminetetraacetate and 0.1% sodium azide, antibody-reactive insulin secreted in the culture supernatant was quantified by enzyme immunoassay or radio immunoassay. The insulin level was indicated as the amount of human insulin (ng/mL). The results are indicated as averages (avg) for 3 to 4 samples with standard error values (se).

Table 3

In the presence of 14.5 mM glucose			
Drug concentration (æM)	Compound No.	Amount of secreted insulin (ng/ml)	
		avg	se
0		148.4	4.8
1.0	12	227.8	3.9
1.0	16	229.5	14.3
1.0	18	181.9	1.4
1.0	19	223.8	3.1
1.0	20	235.9	2.9
1.0	23	249.9	21.3
1.0	31	220.5	2.5
1.0	35	232.4	10.6
1.0	37	202.8	8.0
1.0	43	199.1	3.6
1.0	50	209.5	1.5
1.0	54	216.1	6.6
1.0	65	207.8	2.4
1.0	AY4166	195.1	4.3
0.1	Glibenclamide	177.8	3.3

[0157] As shown in Table 3, it was revealed that the compounds of the present invention had insulin secretion enhancing action. Whilst, in the presence of glucose at a low concentration (5 mM), these compounds did not show marked secretion enhancing action even at a 10 times higher concentration. Glibenclamide (Pharmacotherapy, vol. 5, p.43, 1985) and AY-4166 (Yakuri To Rinsho [Pharmacology and Clinic], vol. 7, p.121, 1997) used as controls for comparison showed marked secretion enhancing action even at a low glucose concentration.

Test Example 3: Insulin secretion enhancing action (glucose-loading)

[0158] Wistar male rats (body weight: about 250 g) starved for 18 hours were anesthetized by intraperitoneal administration of sodium pentobarbital (50 mg/kg), and fixed at the dorsal position. Cannulas were inserted into the right common carotid artery for blood collection and the left femoral vein for administration of drug and glucose, respectively. To these rats were intravenously administered 500 U/kg of heparin to prevent blood coagulation. After about 0.2 ml of blood was collected, a drug was intravenously administered. Two minutes after the administration of the drug, blood was collected, and then glucose (10 mg/kg) was immediately administered intravenously. Then, blood was collected every 20 seconds for 2 minutes. Insulin concentration in plasma obtained by centrifugation was measured by radioimmunoassay. From an insulin concentration which increased after the glucose-loading, its area was calculated and considered the amount of secreted insulin. The insulin level was indicated as the amount of rat insulin (ng/mL). The results are shown in Table 4. As clearly shown in Table 4, the compounds of the present

invention showed significant insulin secretion enhancing action after the glucose-loading (condition of hyperglycemia). On the other hand, they exhibited no secretion enhancing action at a low glucose concentration (normal condition).

Table 4

Test compound	Control		Test compound-administered group				Significant difference
	n	Area	Dose (mg/kg, iv)	n	Area	% Ratio based on control	
16	6	7.02 \pm 0.97	3	6	17.28 \pm 1.35	246 \pm 19	***
19	6	4.61 \pm 3.17	1	6	13.41 \pm 1.95	291 \pm 42	*
35	6	8.83 \pm 2.17	1	6	15.69 \pm 2.04	178 \pm 23	*
23	6	5.19 \pm 0.40	1	6	11.85 \pm 2.58	228 \pm 50	*
31	6	5.19 \pm 0.40	1	6	9.00 \pm 0.84	173 \pm 16	**

Significant difference:

* P < 0.05,

** P < 0.01,

*** P < 0.001 (student's t-test or Aspin-Welch test)

Test Example 4: Hyperglycemia suppression action after glucose-loading

[0159] Wistar male rats (body weight: about 250 g) were anesthetized by intraperitoneal administration of sodium pentobarbital (50 mg/kg), and a cannula was fixed in the left common carotid artery. These rats were used for the experiment after starvation for 18 hours. A test compound was orally administered 1 hour before oral administration of glucose (2 g/kg). Jugular venous blood was collected before the administration of test compound, before the glucose-loading, 5, 10, 15, 30, 45 and 60 minutes after the glucose-loading, and plasma glucose concentration was measured (enzymatic method). The insulin level was indicated as amount of rat insulin (ng/mL). The results are shown in Table 5. As clearly shown in Table 5, the compound of the present invention showed hyperglycemia suppression action after glucose-loading (condition of hyperglycemia). On the other hand, they exhibited no hypoglycemic action under the normal condition.

Table 5

Plasma glucose concentration (mg/dl)						
Test compound	Dose (mg/kg, po)	n	-1 minute	0 minute	5 minutes	10 minutes
Control	-	5	114 \pm 5.0	116 \pm 7.8	152 \pm 6.9	204 \pm 10.7
Compound 16	30	5	114 \pm 4.5	102 \pm 4.6	147 \pm 5.7	183 \pm 9.1
Test compound	Dose (mg/kg, po)	n	15 minutes	30 minutes	45 minutes	60 minutes
Control	-	5	231 \pm 11.8	201 \pm 12.5	189 \pm 9.9	189 \pm 3.2
Compound 16	30	5	186 \pm 8.7*	167 \pm 3.3*	165 \pm 4.8	178 \pm 4.7

Significant difference:

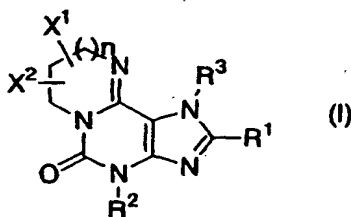
* P < 0.05 (student's t-test or Aspin-Weich test)

Industrial Applicability

[0160] The medicament of the present invention has insulin secretion enhancing action and hypoglycemic action. In particular, the medicament is characterized to have superior hypoglycemic action under a hyperglycemic condition, whilst have no hypoglycemic action under a normal blood glucose level. Therefore, the medicament of the present invention is extremely useful to safely and effectively perform the therapeutic treatment of diabetes and the prophylactic and/or therapeutic treatment of complications of diabetes.

Claims

1. A medicament for the therapeutic treatment of diabetes which comprises as an active ingredient a compound represented by the general formula (I):



- wherein R¹ represents a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aryl group, or a substituted or unsubstituted aromatic heterocyclic group, R² represents a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aralkyl group, a substituted or unsubstituted aryl group, or a substituted or unsubstituted aromatic heterocyclic group, R³ represents a hydrogen atom, a lower alkyl group, or a substituted or unsubstituted aralkyl group, X¹ and X² independently represent a hydrogen atom, a lower alkyl group, a substituted or unsubstituted aralkyl group, or a substituted or unsubstituted aryl group, and the symbol "n" represents an integer of from 0 to 3, or a physiologically acceptable salt thereof.
2. The medicament according to Claim 1, wherein R¹ represents a lower alkyl group, a substituted or unsubstituted aryl group, or a substituted or unsubstituted aromatic heterocyclic group, R² represents a lower alkyl group or a substituted or unsubstituted aryl group, R³ represents a hydrogen atom or a substituted or unsubstituted aralkyl group, X¹ represents a hydrogen atom, a lower alkyl group, or a substituted or unsubstituted aralkyl group, X² represents a hydrogen atom, and the symbol "n" represents 0.
 3. The medicament according to Claim 1, wherein R¹ represents a n-propyl group, a n-butyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, a cyclopropylmethyl group, a cyclopentylmethyl group, an isopropyl group, or a tert-butyl group, R² represents a n-propyl group or an ethyl group, R³ represents a hydrogen atom, X¹ represents a lower alkyl group or a substituted or unsubstituted aralkyl group, X² represents a hydrogen atom, and the symbol "n" represents 0.
 4. A medicament for the prophylactic and/or therapeutic treatment of a complication of diabetes, which comprises a compound represented by the general formula (I) according to Claim 1 or a physiologically acceptable salt thereof as an active ingredient.
 5. A hypoglycemic agent comprising a compound represented by the general formula (I) according to Claim 1 or a physiologically acceptable salt thereof as an active ingredient.
 6. An insulin secretion enhancer comprising a compound represented by the general formula (I) according to Claim 1 or a physiologically acceptable salt thereof as an active ingredient.
 7. Use of a compound represented by the general formula (I) according to Claim 1 or a physiologically acceptable salt thereof for a manufacture of the medicament for the therapeutic treatment of diabetes according to Claim 1.
 8. A method for the therapeutic treatment of diabetes, which comprises the step of administering a therapeutically effective amount of the compound represented by the general formula (I) according to Claim 1 or a physiologically acceptable salt thereof to a mammal including human.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP99/03583

A. CLASSIFICATION OF SUBJECT MATTER
Int.Cl⁶ A61K31/505

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Int.Cl⁶ A61K31/505, C07D487/14

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
CA (STN), REGISTRY (STN)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	JP, 3-204880, A2 (Kyowa Hakko Kogyo Co., Ltd.), 6 September, 1991 (06. 09. 91) & EP, 423805, A2 & US, 5270316, A	1-7
A	JP, 10-158267, A2 (K.K. Medei Sci Puraningu), 16 June, 1998 (16. 06. 98) (Family: none)	1-7
A	D.L. TEMPLE, Jr., J.P. YEVICH, J.D. CATT, D. OWENS, C. HANNING, R.R. CONVINGTON, R.J. SEIDEHAMEL, K.W. DUNGAN, "Substituted 6,7-Dihydroimidazo[1,2- a]purin-9(4H)-ones", Journal of Medicinal Chemistry, 1980, Vol. 23, No. 11, p.1188-1198	1-7
A	FUMIO SUZUKI, JUNICHI SHIMADA, SHIZUKO SHIOZAKI, SHUNJI ICHIKAWA, AKIO ISHII, JOJI NAKAMURA, HIROMI NONAKA, HIROYUKI KOBAYASHI, EIICHI FUSE, "Adenosine A1 Antagonists. 3. Structure-Activity Relationships on Amelioration Against Scopolamine- or N6-((R)-Phenylisopropyl)adenosine-Induced Cognitive Disturbance", Journal of Medicinal Chemistry, 1993, Vol. 36, No. 17, p.2508-2518	1-7

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:
 "A" document defining the general state of the art which is not
 considered to be of particular relevance
 "E" earlier document but published on or after the international filing date
 "L" document which may throw doubts on priority claim(s) or which is
 cited to establish the publication date of another citation or other
 special reason (as specified)
 "O" document referring to an oral disclosure, use, exhibition or other
 means
 "P" document published prior to the international filing date but later than
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 date and not in conflict with the application but cited to understand
 the principle or theory underlying the invention
 "X" document of particular relevance; the claimed invention cannot be
 considered novel or cannot be considered to involve an inventive step
 when the document is taken alone
 "Y" document of particular relevance; the claimed invention cannot be
 considered to involve an inventive step when the document is
 combined with one or more other such documents, such combination
 being obvious to a person skilled in the art
 "&" document member of the same patent family

Date of the actual completion of the international search
16 July, 1999 (16. 07. 99)Date of mailing of the international search report
27 July, 1999 (27. 07. 99)Name and mailing address of the ISA/
Japanese Patent Office

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INTERNATIONAL SEARCH REPORT

International application No.
PCT/JP99/03583

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO, 98/15555, A1 (Kyowa Hakko Kogyo Co., Ltd.), 16 April, 1998 (16. 04. 98) & EP, 884318, A1	1-7

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INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP99/03583

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 8
because they relate to subject matter not required to be searched by this Authority, namely:
It pertains to methods for treatment of the human body by therapy.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

- Remark on Protest**
- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

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